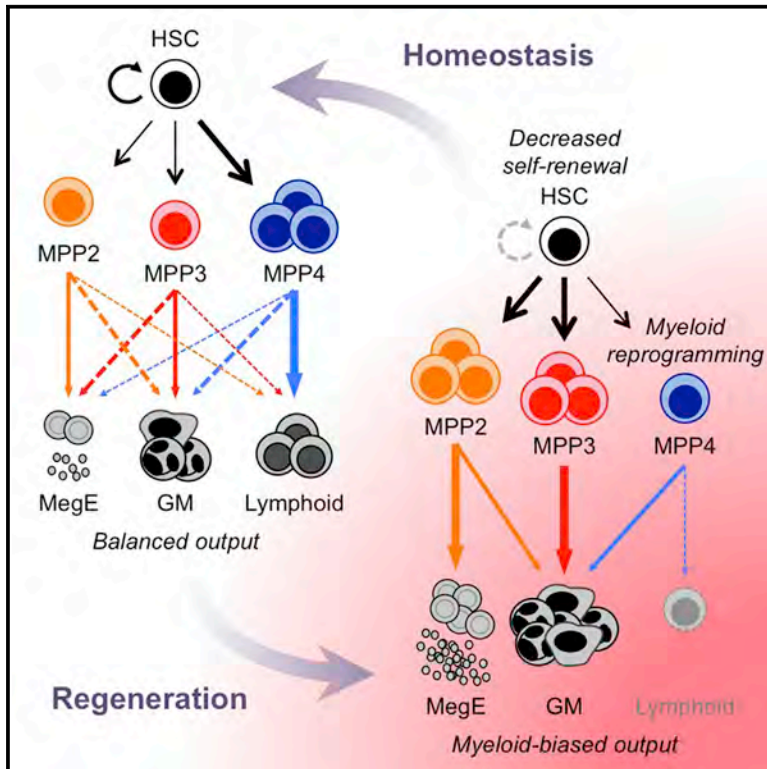


Functionally Distinct Subsets of Lineage-Biased Multipotent Progenitors Control Blood Production in Normal and Regenerative Conditions

Graphical Abstract



Authors

Eric M. Pietras, Damien Reynaud, Yoon-A. Kang, ..., Joshua M. Stuart, Berthold Göttgens, Emmanuelle Passegué

Correspondence

passeguée@stemcell.ucsf.edu

In Brief

Pietras et al. show that HSCs produce functionally distinct lineage-biased MPPs that work together to adapt blood production to hematopoietic demands. Two rare subsets of myeloid-biased MPPs are important for maintaining blood homeostasis at steady state and serve as a potent source of myeloid amplification in regenerative conditions.

Highlights

- Blood production is mediated by functionally distinct lineage-biased MPPs
- MPP subsets are independently produced by HSCs
- Myeloid-biased MPPs serve as a potent source of myeloid amplification
- Regeneration occurs at the expense of HSC engraftment and self-renewal activity

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Functionally Distinct Subsets of Lineage-Biased Multipotent Progenitors Control Blood Production in Normal and Regenerative Conditions

Eric M. Pietras,^{1,5} Damien Reynaud,^{1,5,6} Yoon-A. Kang,¹ Daniel Carlin,² Fernando J. Calero-Nieto,³ Andrew D. Leavitt,⁴ Joshua M. Stuart,² Berthold Göttgens,³ and Emmanuelle Passegué^{1,*}

¹The Eli and Edythe Broad Center of Regeneration Medicine and Stem Cell Research, Division of Hematology/Oncology, Department of Medicine, University of California San Francisco, San Francisco, CA 94143, USA

²Department of Biomolecular Engineering and Center for Biomolecular Science and Engineering, University of California Santa Cruz, Santa Cruz, CA 94720, USA

³Cambridge University Department of Haematology, Cambridge Institute for Medical Research and Wellcome Trust and MRC Cambridge Stem Cell Institute, Hills Road, Cambridge CB2 0XY, UK

⁴Departments of Medicine and Laboratory Medicine, University of California San Francisco, San Francisco, CA 94143, USA

⁵Co-first author

⁶Present address: Division of Experimental Hematology and Cancer Biology, Cincinnati Children's Hospital Medical Center, Cincinnati, OH 45229, USA

*Correspondence: passeguée@stemcell.ucsf.edu

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SUMMARY

Despite great advances in understanding the mechanisms underlying blood production, lineage specification at the level of multipotent progenitors (MPPs) remains poorly understood. Here, we show that MPP2 and MPP3 are distinct myeloid-biased MPP subsets that work together with lymphoid-primed MPP4 cells to control blood production. We find that all MPPs are produced in parallel by hematopoietic stem cells (HSCs), but with different kinetics and at variable levels depending on hematopoietic demands. We also show that the normally rare myeloid-biased MPPs are transiently overproduced by HSCs in regenerating conditions, hence supporting myeloid amplification to rebuild the hematopoietic system. This shift is accompanied by a reduction in self-renewal activity in regenerating HSCs and reprogramming of MPP4 fate toward the myeloid lineage. Our results support a dynamic model of blood development in which HSCs convey lineage specification through independent production of distinct lineage-biased MPP subsets that, in turn, support lineage expansion and differentiation.

INTRODUCTION

Blood production is a highly regulated process that tailors the output of the myeloid and lymphoid lineages based on hematopoietic demands and the needs of the organism (Ema et al., 2014). Blood development starts with rare self-renewing hematopoietic stem cells (HSCs), which produce a series of increasingly more abundant and lineage-committed progenitor cells, ultimately giving rise to all types of mature blood cells.

Although the overall structure of the blood system and its hierarchical nature is well established, many questions still remain regarding how HSCs specify lineage fate in non-self-renewing multipotent progenitors (MPPs) prior to the generation of lineage-committed progenitors and the separation of the myeloid and lymphoid lineages.

HSCs are defined functionally by their ability to serially engraft transplanted recipients and regenerate the entire blood system. This unique property is used to directly measure HSC self-renewal activity and to identify HSCs based on phenotypic markers. In the mouse, HSCs are found in the Lin[−]/Sca-1⁺/c-Kit⁺ (LSK) fraction of the bone marrow (BM) and are usually defined as CD150⁺/CD48[−] LSK cells (Kiel et al., 2005), although other surface markers can be used to enrich for more quiescent and/or functionally distinct subsets including Flk2, CD34, EPCR, rhodamine, the other SLAM markers CD229 and CD244, and CD41 (Wilson et al., 2008; Kent et al., 2009; Oguro et al., 2013; Yamamoto et al., 2013; Miyawaki et al., 2015). Transplantation experiments have shown that markers enriching for the most quiescent and metabolically inert HSC subsets will directly favor engraftment and self-renewal activity (Pietras et al., 2011; Kohli and Passegué, 2014). Single cell transplantation experiments have further demonstrated that even HSCs with identical surface phenotypes are heterogeneous in their engraftment behaviors, with different stabilities over time and variable degrees of myeloid versus lymphoid lineage output (Dykstra et al., 2007; Yamamoto et al., 2013). Several models are currently proposed to explain this heterogeneity, including the sequential loss of lineage potential in differentiating HSCs (Adolfsson et al., 2005), the existence of long-lived myeloid bypass pathways (Yamamoto et al., 2013), and the presence of functionally distinct clones of lineage-biased HSCs with different biological activities and hard-wired lineage potentials (Muller-Sieburg et al., 2004; Dykstra et al., 2007). However, it remains unclear whether the behavior of transplanted HSCs accurately reflects steady-state hematopoiesis and HSC function in native conditions. In fact, it is possible that the distinct

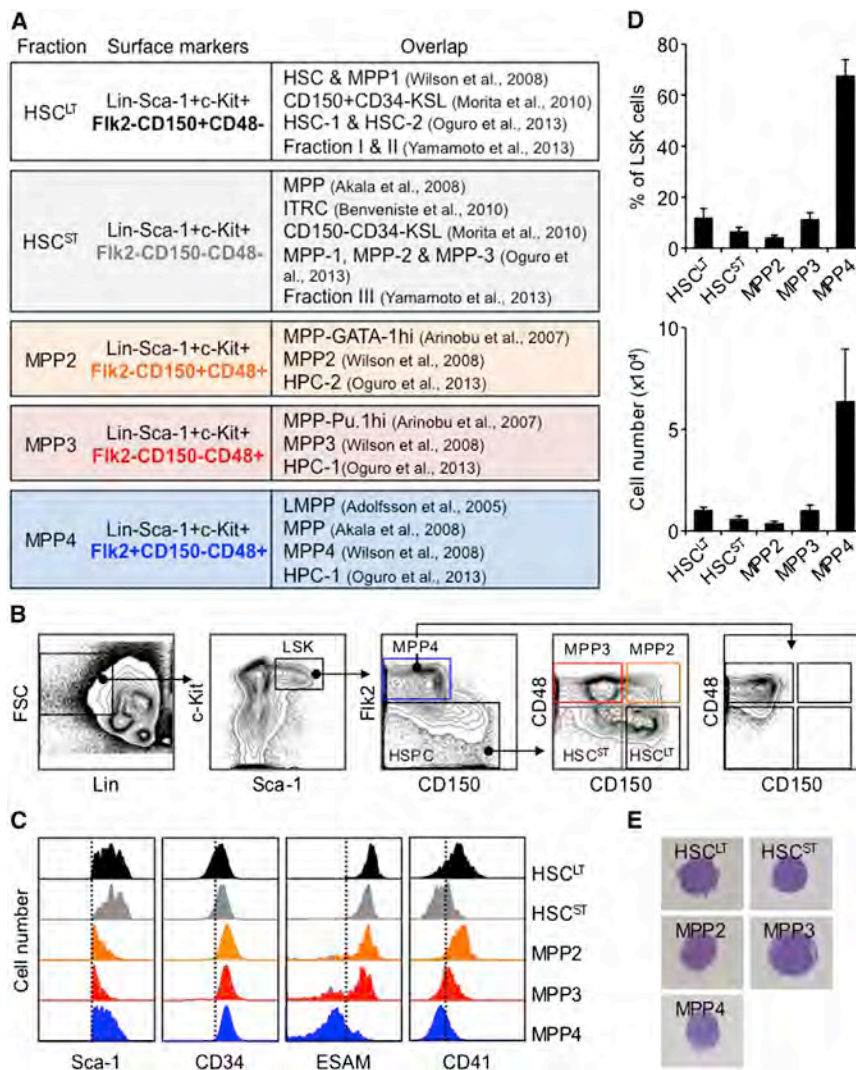


Figure 1. Reinvestigating MPP Subsets

(A) Table showing overlap of MPP subsets with previously published definitions.

(B) Representative gating strategy used to identify and isolate HSC^{LT}, HSCST, MPP2, MPP3, and MPP4 based on expression of Fik2, CD48, and CD150 in BM LSK.

(C) Representative histograms of Sca-1, CD34, ESAM, and CD41 expression in the indicated LSK subsets.

(D) Average percentage in BM LSK and absolute numbers of each population (8 mice/group).

(E) Wright-Giemsa staining of the indicated LSK subsets.

Results are expressed as mean \pm SD.

et al., 2006; Boyer et al., 2011; Buza-Vidas et al., 2011). In fact, the top 25% of the most highly expressing Fik2⁺ LSK cells have been called lymphoid-primed MPPs or LMPPs (Adolfsson et al., 2005). Recently, two other MPP subsets have been described in the Fik2⁻ LSK fraction and termed MPP2 and MPP3, with Fik2⁺ LSK cells re-named MPP4 (Wilson et al., 2008). Although preliminary investigations suggest that MPP2 and MPP3 have myeloid-biased outputs (Cabezas-Wallscheid et al., 2014), little is known about their biological function. Here, we directly compared the function of MPP2, MPP3, and MPP4 in blood production at steady state and in regenerating conditions following HSC transplantation. We propose a model wherein HSCs produce in parallel distinct subsets of lineage-biased MPPs, which together coordinate the output of the

behaviors exhibited by single transplanted HSCs represent reversible activities that span a continuum of surface markers and activation states. The identification of MPP1 as a metabolically active subset of HSCs directly supports this idea (Cabezas-Wallscheid et al., 2014). Moreover, two exciting lineage tracking studies marking HSCs in their native environment using either sleeping beauty transposons (Sun et al., 2014) or *Tie2-Cre* endogenous labeling (Busch et al., 2015) have recently shown a limited contribution of HSCs to steady-state hematopoiesis and, conversely, a major role for MPPs and lineage-committed progenitors to ongoing blood production.

MPPs are currently a poorly defined hematopoietic compartment, and the term itself is used rather indiscriminately to refer to cells within the LSK fraction that have limited to no engraftment ability in transplantation experiments. The best-characterized and most abundant MPP subset is defined as Fik2⁺ LSK cells and is now considered as a fully multipotent but lineage-biased population, with low megakaryocyte/erythroid (MegE) and high lymphoid potentials (Adolfsson et al., 2005; Forsberg

myeloid and lymphoid lineages in response to hematopoietic demands.

RESULTS

Lineage-Biased MPP Subsets

Many names and phenotypic definitions are currently used to describe the spectrum of MPP subsets present in the mouse LSK BM compartment (Figure 1A). One of the most broadly applicable schemes separates the most quiescent HSCs (CD34⁻/Fik2⁻/CD150⁺/CD48⁻ LSK) from the more metabolically active MPP1 (CD34⁺/Fik2⁻/CD150⁺/CD48⁻ LSK) and divides MPPs into three further distinct subsets: MPP2 (Fik2⁻/CD150⁺/CD48⁺ LSK), MPP3 (Fik2⁻/CD150⁻/CD48⁺ LSK), and MPP4 (Fik2⁺/CD150⁻/CD48⁺ LSK) (Figure 1B) (Wilson et al., 2008; Cabezas-Wallscheid et al., 2014). Importantly, these populations overlap with other MPP definitions based on reporter gene combinations and different surface markers (Arinobu et al., 2007; Akala et al., 2008; Oguro et al., 2013). Because our focus was on MPP biology, we did not use CD34 separation and referred

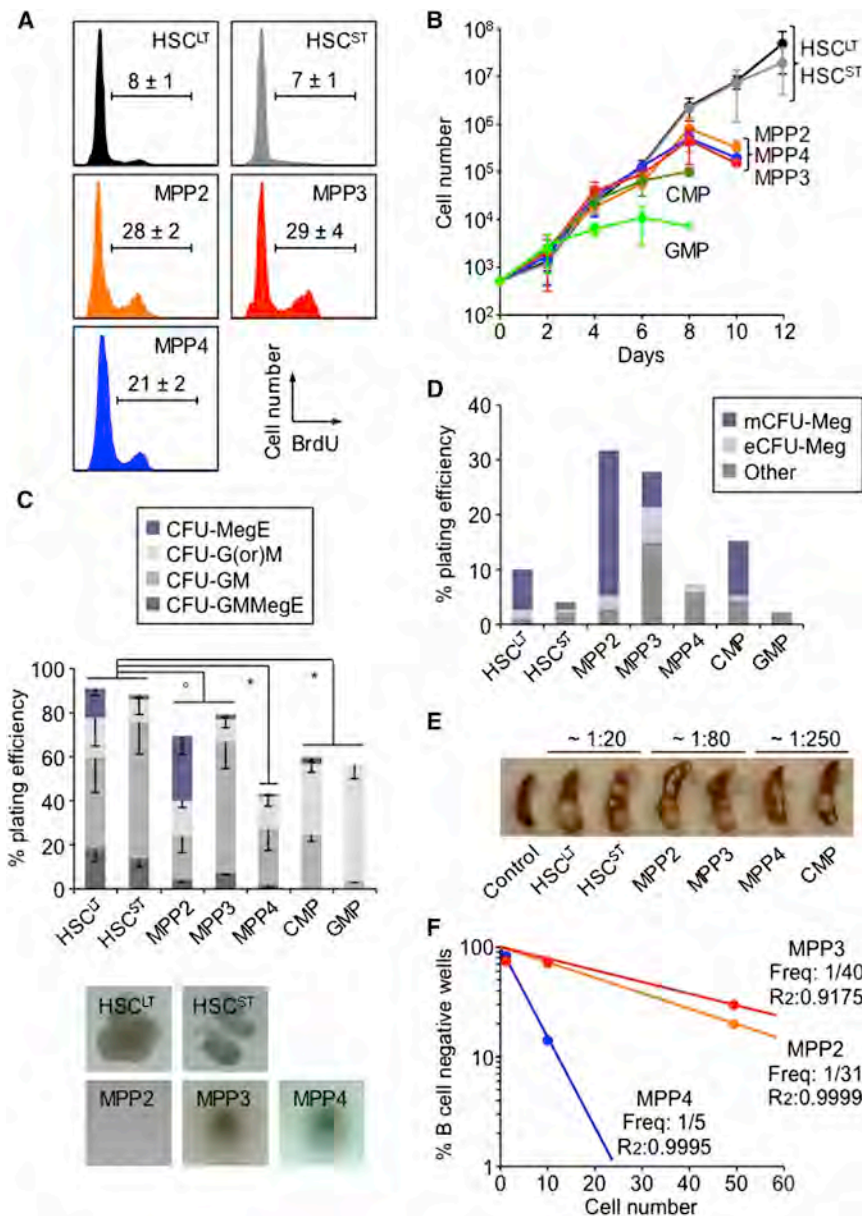


Figure 2. Coexistence of Functionally Distinct MPP Subsets

(A) Proliferation rates. Cells were pulsed for 1 hr with BrdU before analysis ($n \geq 3$).

(B) Expansion in liquid culture. Cells were counted every other day ($n = 2$).

(C) Methylcellulose clonogenic assays and pictures of representative colonies. Single cells were used to measure plating efficiency and colony-forming unit (CFU) activity ($n \geq 3$).

(D) Meg differentiation potential in collagen-based MegaCult assays ($n = 2$). mCFU-Meg: small mature colony of ≤ 6 Meg; eCFU-Meg: large early colony of ≥ 6 Meg.

(E) CFU-S assays. Representative photograph of spleen colonies obtained after transplantation of the indicated populations in lethally irradiated mice ($n = 2$). CFU-S frequency is given at day 12.

(F) Clonogenic B cell differentiation potential on OP9/IL-7 stromal cells. Single, 10, and 50 cells were grown for 16 days and analyzed by flow cytometry for production of CD19⁺ B cells (10–34 wells/cell dose).

Results are expressed as mean \pm SEM; $\circ p < 0.05$, $\ast p < 0.001$.

See also Figure S1.

to the combination of HSC and MPP1 as HSC^{LT}. We also included an additional sub-population of LSK cells defined as CD34⁺/Flk2[−]/CD150[−]/CD48[−] LSK (Figure 1B) that we named short-term HSCs (HSCST) and referred to the combination of HSC^{LT} and HSCST as HSCs.

Both MPP2 and MPP3 showed low surface expression of Sca-1 compared to HSCs and MPP4 and, like MPP4, expressed high levels of CD34 (Figure 1C). However, unlike MPP4, they also expressed the HSC markers ESAM (Ooi et al., 2009) and, to various degrees, CD41 (Figure 1C). In steady-state conditions, both MPP2 and MPP3 were as rare as HSCs (Figure 1D), and were morphologically undistinguishable from other LSK populations (Figure 1E). As expected, short-term bromodeoxyuridine (BrdU) incorporation experiments showed higher proliferation rates in MPP2 and MPP3 than in HSCs, which were in the range

of MPP4 (Figure 2A) (Wilson et al., 2008). Liquid culture experiments confirmed similar expansion rates from all three MPP subsets, which distinguished them from the long-lived HSCs and growth-restricted common myeloid progenitors (CMPs) and granulocyte/macrophage progenitors (GMPs) (Figure 2B). Flow cytometry analyses for Mac-1 and FcγR myeloid differentiation markers also demonstrated persistence of granulocyte/macrophage (GM) potential in cultured MPP2 and MPP3 compared to the transient myeloid output observed with MPP4 (Figure S1A). In clonogenic methylcellulose assays, both MPP2 and MPP3 gave rise to all myeloid lineages but with plating efficiency and pattern

of differentiation distinct from HSCs, MPP4, and myeloid progenitors (Figure 2C). Strikingly, MPP3 showed a dominant GM output, whereas MPP2 displayed extensive Meg potential, which was confirmed in collagen-based MegaCult assays (Figure 2D). Spleen colony-forming unit (CFU-S) assays also indicated similarly strong erythroid potential from both MPP2 and MPP3, which was intermediate to that of HSCs and MPP4/CMPs (Figure 2E). When cultured on OP9/IL-7 to assess B cell potential, both MPP2 and MPP3 generated CD19⁺ B cells, but with kinetics that again were intermediary between HSC^{LT} and MPP4 (Figure S1B). Furthermore, single cell analyses revealed ~8-fold lower B cell clonogenic potential in MPP2 and MPP3 compared to MPP4, and a unique ability to produce B/myeloid mix colonies (Figures 2F and S1C). Similarly, when cultured on OP9-DL1 stromal cells for T cell potential, both MPP2 and MPP3 produced

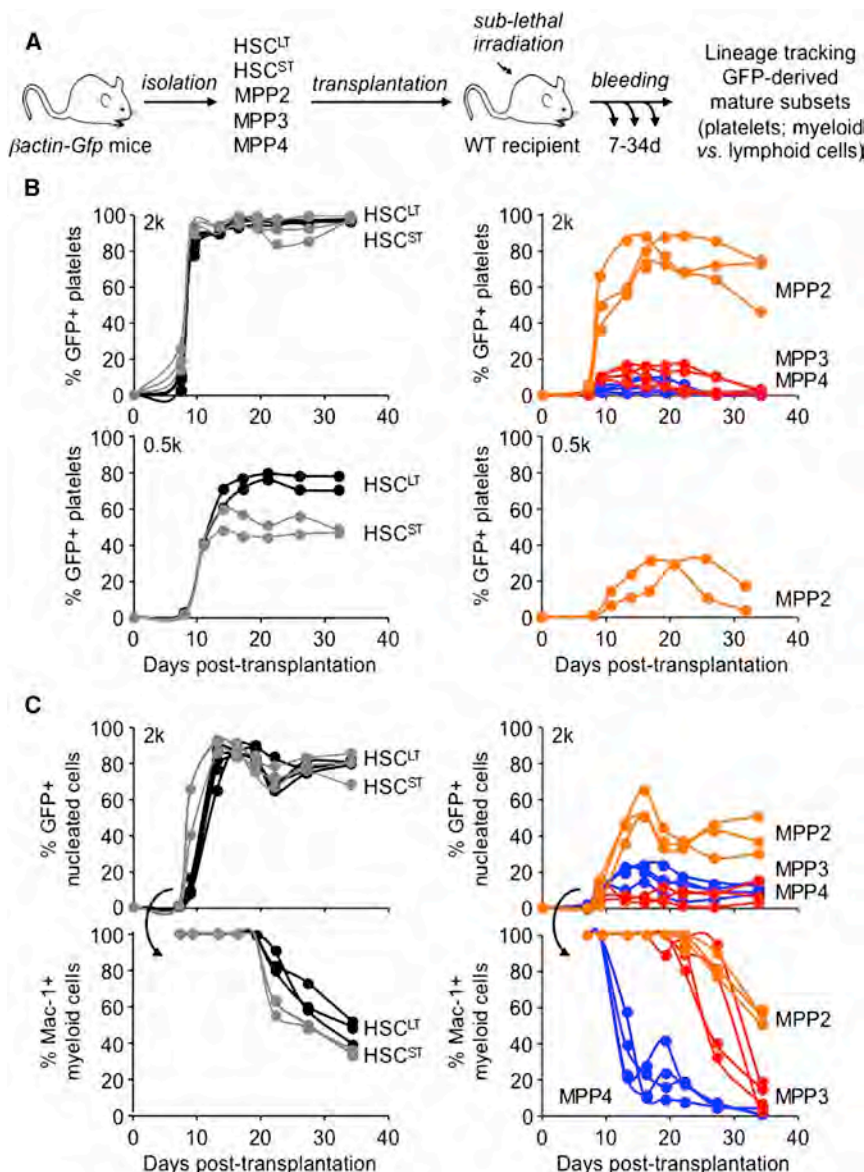


Figure 3. Specific Lineage Biases in MPP Subsets

(A) Experimental scheme for the in vivo lineage tracking experiments. GFP⁺ populations were transplanted into sub-lethally irradiated recipients and followed over time for their reconstitution activity and lineage potential in PB.

(B) Platelet chimerism following transplantation of 2,000 (upper graphs) or 500 (lower graphs) cells of the indicated donor GFP⁺ population. Each line represents individual mice (2-4 mice/group).

(C) Nucleated cell chimerism (upper graphs) and percent of donor-derived Mac1⁺ myeloid cells (lower graphs) following transplantation of 2,000 cells of the indicated donor GFP⁺ population. Each line represents individual mice (2-4 mice/group).

See also Figure S2.

lethally irradiated recipients and followed their multilineage output in peripheral blood (PB) as we have previously done (Forsberg et al., 2006) (Figure 3A). These lineage-tracking experiments confirmed that MPP2 and MPP3 were both devoid of self-renewal potential (Santaguida et al., 2009; Cabezas-Wallscheid et al., 2014) because, like MPP4, they only exhibited short-term myeloid reconstitution ability (≤ 1 months) in contrast to the sustained potential displayed by HSCs (Figure 3B). Interestingly, differences in HSC^{LT} and HSCST reconstitution activity were only revealed by transplanting a low number of cells (Figure 3B) and tracking the persistence of myeloid output for up to 4 months (Figure S2A). In fact, HSCST behaved similarly to the previously described intermediate-term HSCs (ITRC) (Benveniste et al., 2010). Importantly, these lineage-tracking ex-

periments directly showed the multipotent nature of MPP2 and MPP3, and their ability to generate low levels of B and T cells in vivo (Figure S2B). They also highlighted the considerable but transient ability of MPP2 to produce platelets (Figures 3B and S2C), and the extensive GM potential of both MPP2 and MPP3, which were able to maintain production of mature myeloid cells for 2 weeks longer than MPP4 (Figure 3C). Taken together, these in vivo experiments confirm that MPP2 and MPP3 are two myeloid-biased MPP subsets that are functionally distinct from the lymphoid-primed MPP4.

Functional Validation

To further demonstrate the intrinsic myeloid bias of these two MPP subsets, we transplanted HSC^{LT}, HSCST, MPP2, MPP3, and MPP4 isolated from β -actin-Gfp mice into sub-

lethally irradiated recipients and followed their multilineage output in peripheral blood (PB) as we have previously done (Forsberg et al., 2006) (Figure 3A). These lineage-tracking experiments confirmed that MPP2 and MPP3 were both devoid of self-renewal potential (Santaguida et al., 2009; Cabezas-Wallscheid et al., 2014) because, like MPP4, they only exhibited short-term myeloid reconstitution ability (≤ 1 months) in contrast to the sustained potential displayed by HSCs (Figure 3B). Interestingly, differences in HSC^{LT} and HSCST reconstitution activity were only revealed by transplanting a low number of cells (Figure 3B) and tracking the persistence of myeloid output for up to 4 months (Figure S2A). In fact, HSCST behaved similarly to the previously described intermediate-term HSCs (ITRC) (Benveniste et al., 2010). Importantly, these lineage-tracking ex-

Molecular Validation

To understand how MPP2 and MPP3 fit into the hematopoietic continuum, we conducted genome-wide microarray analyses. Hierarchical clustering based on gene signatures derived from the 1,000 most highly differentially expressed genes in HSC^{LT} and GM lineage-committed cells (Figures S3A and S3B;

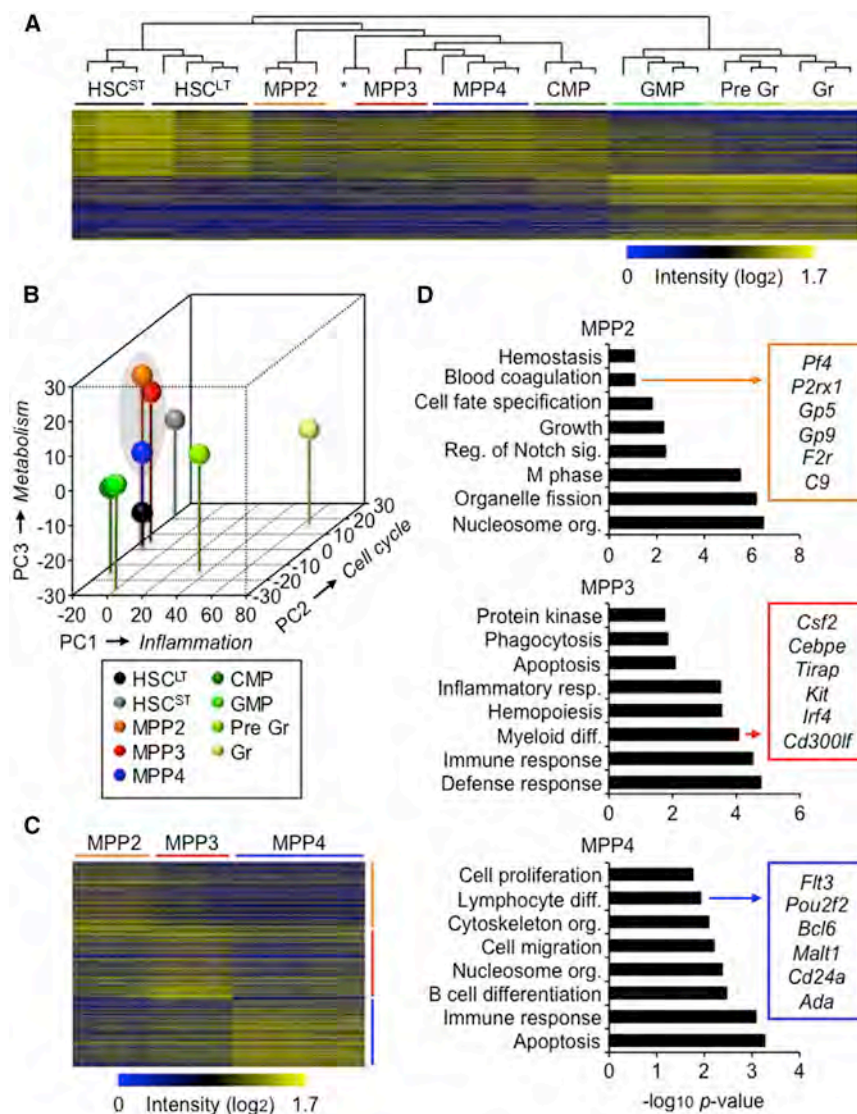


Figure 4. Molecular Biases in MPP Subsets

(A) Hierarchical clustering analysis based on the 1,000 most highly expressed genes in HSC^{LT} and GM lineage-committed cells (GMP, Gr precursors: pre Gr; Gr). *Indicates one MPP4 sample clustering independently.

(B) Principal component (PC) analysis of the microarray results shown in (A). Axis labels indicate the primary gene signature driving each PC separation.

(C) Individual gene signatures representing the 1,000 most highly expressed genes in MPP2, MPP3, and MPP4 relative to the other two populations.

(D) Gene ontology (GO) analyses and examples of the gene signatures shown in (C).

See also Figure S3 and Tables S1, S2, S3, and S4.

active MegE transcription program, with high expression levels of *Gfi1b*, *Gata1*, and *Fog1* (Figure S3D). These distinct gene expression signatures allowed recognition of MPP2, MPP3, and MPP4 in other published gene expression datasets (Table S4). Taken together, these molecular studies corroborate our functional analyses by positioning the rare myeloid-biased MPP2 and MPP3 together with the more abundant lymphoid-primed MPP4 in a multipotent compartment located downstream of HSCs and upstream of the already lineage-committed myeloid and lymphoid progenitors.

Hierarchical Organization

To confirm the hierarchy between MPP2, MPP3, and MPP4, we first performed short-term in vitro differentiation experiments, and followed changes in early

stem and progenitor surface marker expression by flow cytometry (Figures 5A and S4A). Whereas robust c-Kit and Sca-1 expression allowed tracking of LSK differentiation, the lack of Flk2 induction in vitro or its quick downregulation following exposure to Flt-3 ligand did not permit the separation of newly generated MPP3 and MPP4. In both HSC^{LT} and HSCST cultures, we observed a persistent LSK compartment, which first produced MPP2 and then MPP3/4 (Figure 5A). In contrast, in MPP cultures, the LSK compartment quickly differentiated into c-Kit⁺/Sca-1⁺ myeloid progenitors and c-Kit⁺/Sca-1⁺ mature myeloid cells, with the few remaining LSK cells preserving their initial identity despite some fluctuation in marker expression (Figure S4A). Although these in vitro results establish that MPP2 is the first MPP subset produced by differentiating HSCs, they do not distinguish whether MPP2 then make MPP3/4, or whether the other MPP subsets are directly generated from HSCs but with slower kinetics. To further clarify this relationship, we took advantage of mice lacking the thrombopoietin receptor (*Mpl*) gene (Gurney et al., 1994)

Table S1), divided the hematopoietic hierarchy into three main groups: (1) highly immature HSC^{LT} and HSCST; (2) multipotent progenitors including MPP2, MPP3, MPP4, and early myeloid-committed CMPs; and (3) GM lineage-committed cells including GMPs, Gr precursors (Pre Gr), and Gr (Figure 4A). Interestingly, principal component (PC) and gene ontology (GO) analyses identified inflammation genes as main drivers of PC1, which set apart the entire GM-committed myeloid differentiation axis, whereas cell-cycle genes drove PC2, which segregated HSCST from HSC^{LT}, and metabolism genes PC3, which separated MPP2 and MPP3 from MPP4 (Figures 4B and S3C; Table S2). Moreover, GO analyses performed on gene signatures representing the 1,000 most highly differentially expressed genes in each MPP relative to the other two underscored their unique molecular features (Figures 4C and 4D; Table S3). As expected, the MPP4 signature was enriched in lymphoid differentiation genes, whereas the MPP3 signature was enriched for myeloid differentiation genes and MPP2 signature for genes related to hemostasis. Consistently, MPP2 showed features of an

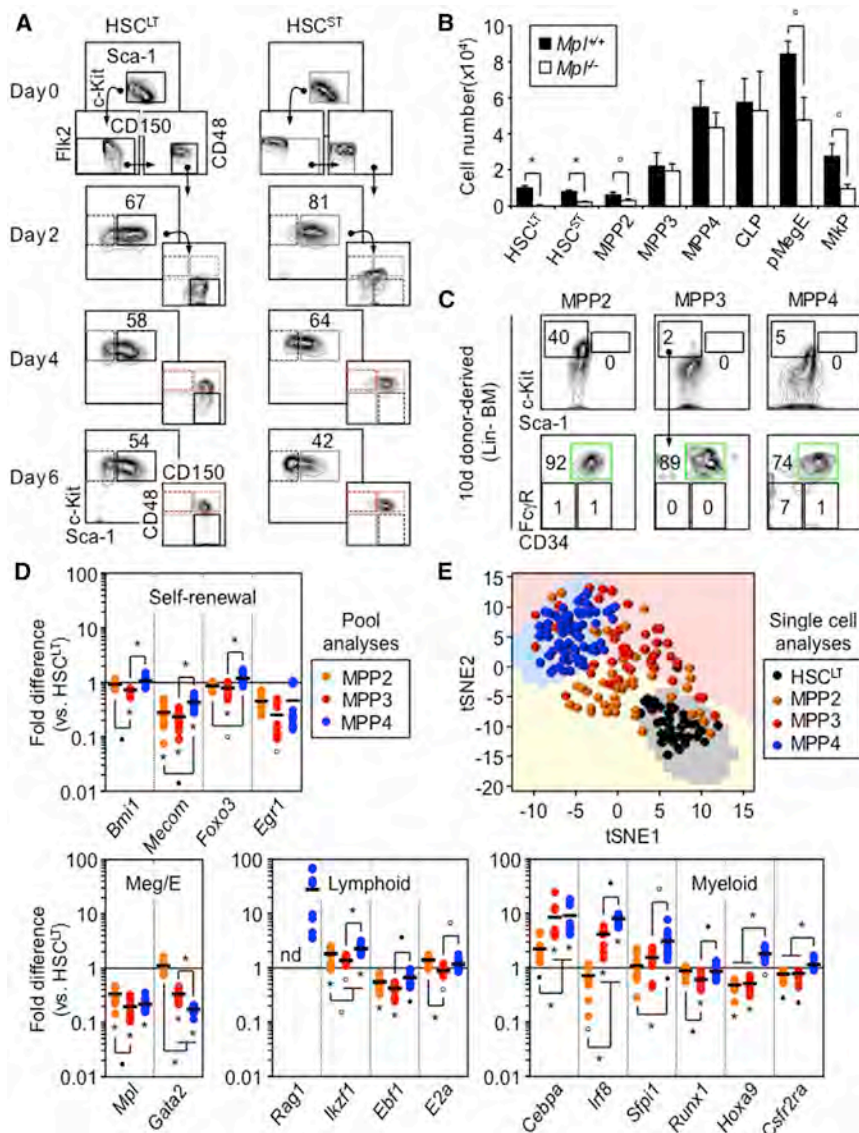


Figure 5. Hierarchical Organization and Molecular Priming

(A) Differentiation in vitro. Representative FACS plots showing HSC^{LT} and HSCST differentiation kinetics in myeloid conditions (n = 2).

(B) Size of the indicated BM populations in *Mpl*^{-/-} and littermate control mice (3 mice/group).

(C) Differentiation in vivo. Representative FACS plots of LSK and myeloid progenitor output 10 days following transplantation of 5,000 cells of the indicated donor population (3 mice/group).

(D) Fluidigm gene expression analyses at steady state. Results are expressed as mean (bar) and individual fold differences compared to HSC^{LT} (8–12 pools of 11 cells/population; nd, not detectable).

(E) t-distributed stochastic neighbor embedding (tSNE) analysis of Fluidigm gene expression data acquired from single cells (30–58 cells/population). Results are expressed as mean ± SD; *p < 0.05, **p < 0.01, ***p < 0.001.

See also Figures S4 and S5.

robust LSK population within 14 days post-transplantation (Figures 5C and S4C). In contrast, after 10 days, we observed strong production of donor-derived GMPs from all MPP subsets, which persisted 14 days after transplantation for MPP2 and MPP3, but not for MPP4, which, by then, was mostly producing lymphoid-derived progeny (Figure S4D). These results confirm the enhanced production of GMPs by myeloid-biased MPP2 and MPP3 and demonstrate that no MPPs are able to generate other MPPs in vivo. Taken together, these experiments establish a hierarchy where HSCs independently generate all three types of lineage-biased MPPs, but with faster production

to determine how defects in platelet production could affect the genesis of the different MPP subsets (Figure 5B). We confirmed the major decrease in HSC^{LT} and HSCST in *Mpl*^{-/-} mice (Qian et al., 2007) and found a consistent ~2-fold reduction in the numbers of MPP2 and all lineage-committed progenitors with Meg activity including Pre-MegE, MkP, and CMP (Figures 5B and S4B). In sharp contrast, the numbers of MPP3, MPP4, and other progenitors devoid of Meg activity remained unchanged. These results highlight the key contribution of the Meg-biased MPP2 to megakaryopoiesis, and suggest that MPP3 and MPP4 are produced independently from MPP2 by HSC^{LT}. Finally, we transplanted MPP2, MPP3, and MPP4 isolated from wild-type mice into sub-lethally irradiated recipients to evaluate their ability to regenerate other MPP subsets shortly after transplantation (Figure 5C). However, despite injecting high cell numbers, we were unable to detect production of donor-derived LSK cells in BM from any MPP subsets, while in the same conditions HSC^{LT} regenerated a

of Meg-biased MPP2 over GM-biased MPP3 and lymphoid-primed MPP4.

Molecular Priming at Steady State

To gain insights into the molecular pathways controlling the production of these different MPP subsets, we used a custom-made Fluidigm dynamic PCR array platform to analyze the expression of 57 genes in pools of 100 HSC^{LT}, MPP2, MPP3, and MPP4 (Figure 5D). Importantly, we detected the expected changes in expression of the surface markers *Slamf1* (CD150), *Cd48*, and *Flt3* (Flk2) that define the identity of these populations (Figure S5A). Overall, we found a significant reduction in expression of self-renewal genes in MPPs compared to HSC^{LT}, which was more pronounced in MPP3 than in MPP4 (Figure 5D), and was confirmed at the protein level using cells isolated from *Bmi1-eGFP* reporter mice (Figure S5B) (Hosen et al., 2007). We also documented the Meg poising of MPP2, with maintenance of *Gata2* expression to levels similar to HSC^{LT}, and the lymphoid

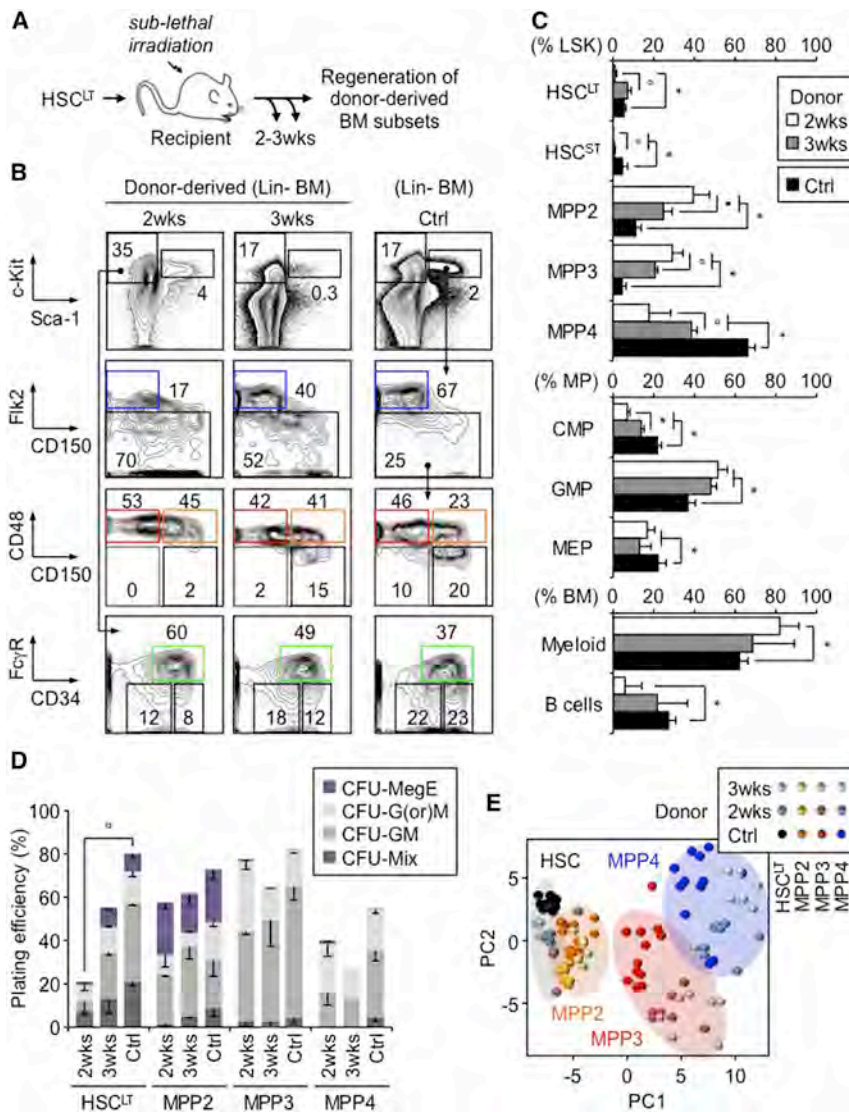


Figure 6. Contribution to Blood Regeneration

(A) Experimental scheme for in vivo blood regeneration experiments. Donor HSC^{LT} were transplanted into sub-lethally irradiated congenic recipients (2,000 HSC^{LT}/mouse) and regeneration of donor BM subsets was followed at 2 and 3 weeks post-transplantation and compared to steady-state control (Ctrl) mice.

(B) Representative FACS plots of regenerating BM subsets.

(C) Frequency of the indicated BM subsets (6–10 mice/group).

(D) Methylocellulose clonogenic assays for the indicated populations (n = 1–4)

(E) PC analysis of Fluidigm gene expression data from the indicated populations (8–12 pools of 100 cells/condition).

Results are expressed as mean ± SD or SEM (D); ○ p < 0.05, ● p < 0.01, *p < 0.001.

See also Figures S5 and S6.

In addition, *Cebpa*, *Irf8*, *Ccne1/2* (cyclin E1 and E2), and *Rad51* emerged as important drivers for both PC1 and PC2 separation, which together with *Slamf1* distinguished MPP2 and MPP3 from each other and separated them from HSC^{LT} and MPP4. Collectively, these analyses demonstrate that, at steady state, all three MPP subsets are part of an actively proliferating continuum of differentiation with molecular priming toward their respective lineage biases and, in the case of the most abundant MPP4, persistence of a significant GM poised alongside its lymphoid priming.

Contribution to Blood Regeneration

To understand how each MPP subset emerges from HSCs and contributes to

priming of MPP4, with induction of *Rag1* and higher expression of *Ikzf1* (Ikaros) compared to HSC^{LT} and other MPP subsets (Figure 5D). In addition, both MPP3 and MPP4 showed clear evidence of GM poising compared to HSC^{LT} and MPP2, with induction of *Cebpa*, *Irf8*, and *Sfp1* (Pu.1) gene expression and protein levels as measured in cells isolated from *Pu.1-eYFP* reporter mice (Figures 5D and S5B). Consistent with their active cell-cycle status, all MPPs displayed higher expression levels of specific G1 to S and G2 to M phase cyclin/CDK complex genes compared to HSC^{LT} and a major downregulation of the quiescence-enforcer *Cdkn1c* (p57) (Figure S5C). All MPPs also showed elevated expression levels of some DNA repair pathway components, especially *Rad51* and *Rpa1* (Figure S5C). Moreover, we performed single cell analyses with the same Fluidigm platform and investigated the 49 most robustly expressed genes using PC analyses and t-distributed stochastic neighbor embedding (tSNE) to pinpoint the key molecular drivers segregating these different populations (Figures 5E and S5D). As expected, *Cd48*, *Slamf1*, and *Flt3* were all important for PC1 separation.

blood regeneration, we injected CD45.2 donor HSC^{LT} into sub-lethally irradiated CD45.1 congenic recipients and followed the production of donor-derived BM populations at 2 and 3 weeks post-transplantation (Figure 6A). Strikingly, at 2 weeks, both MPP2 and MPP3 were massively expanded and accounted for most of the reforming LSK compartment, with HSC^{LT} and HSCST being exceedingly rare and MPP4 just starting to be generated (Figures 6B and 6C). By 3 weeks, this process began to revert, with MPP2 and MPP3 contracting toward their steady-state levels as HSC^{LT} and MPP4 expanded. In addition, at 2 weeks, all donor-derived BM cells were essentially myeloid, whereas by 3 weeks, lymphopoiesis had recovered, with a BM lineage composition becoming similar to steady-state levels (Figures 6C and S5E). Consistently, we observed a transient expansion of GMPs at 2 weeks, with kinetics matching the initial burst of MPP2/MPP3 expansion and myeloid cell production (Figure 6C). These results indicate that regenerating HSCs first produce myeloid-biased MPPs to quickly establish myeloid output, followed by lymphoid-primed MPP4 to rebuild the lymphoid

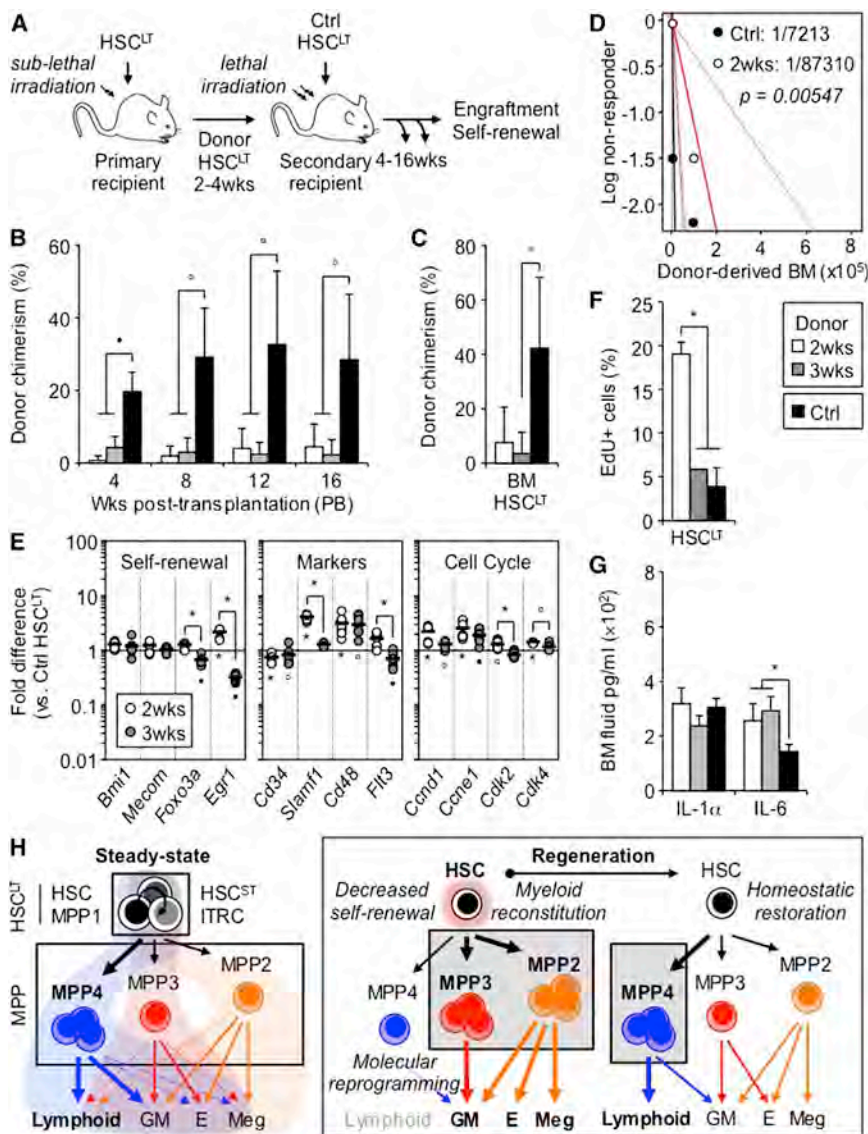


Figure 7. Myeloid-Biased MPPs Are a Transient Compartment of Myeloid Amplification

(A) Experimental scheme for re-transplantation of regenerating HSC^{LT}. Donor CD45.2 HSC^{LT} isolated from primary recipients at 2 and 3 weeks post-transplantation or from Ctrl mice were injected into lethally irradiated secondary CD45.1 recipients (100 HSC^{LT}/mouse) together with 3×10^5 Sca-1-depleted CD45.1 BM cells (4–13 mice/group).

(B) Engraftment over time in PB.

(C) Engraftment in BM HSC^{LT} at 16 weeks post-transplantation.

(D) Limit dilution analyses (LDA) of Ctrl (black) and 2 weeks post-transplantation (red) BM cells. Dotted lines represent confidence intervals and values the estimated HSC frequency.

(E) Fluidigm gene expression analyses of key self-renewal determinant, surface marker, and cell-cycle genes in regenerating HSC^{LT} at 2 and 3 weeks post-transplantation. Results are expressed as mean (bar) and individual fold compared to steady-state Ctrl HSC^{LT} (8–12 pools of 100 cells/condition).

(F) Proliferation rates in mice pulsed for 1 hr with EdU (n = 1–3).

(G) IL-1 and IL-6 levels in BM fluid (5 mice/group).

(H) Revised model of blood production at steady state and in regenerating conditions.

Results are expressed as mean \pm SD or SEM (F); \circ p < 0.05, \bullet p < 0.01, *p < 0.001.

See also Figure S7.

lineages. Functional assessment in methylcellulose showed similar plating efficiency and overall lineage distribution between newly produced MPP subsets at 2 and 3 weeks, compared to steady state (Figure 6D). In contrast, regenerating HSC^{LT} at 2 weeks displayed profoundly impaired differentiation potential in vitro, which was in large part recovered by 3 weeks (Figure 6D).

We also performed Fluidigm gene expression analyses on pools of 100 donor-derived cells re-isolated at 2 and 3 weeks post-transplantation to gain a molecular understanding of this regeneration process (Figures 6E and S6). PC analyses indicated that regenerating HSC^{LT} and newly generated MPP2 preserved their overall molecular identity, whereas MPP3 and MPP4 showed greater dispersion, mainly reflecting more pronounced changes in expression levels of certain lineage commitment genes (Figure 6E). In particular, Meg/E lineage determinants were significantly downregulated in MPP3 and MPP4, while being preserved in both HSC^{LT} and MPP2 (Figure S6). In addition, MPP4 showed changes in both lymphoid lineage determinants, with transient upregulation of

MPP2 and MPP3 displayed limited changes in GM lineage determinants aside from increased expression of *Irf8* at 3 weeks and fluctuations in *Hoxa9* expression. These results show that MPP4 undergo a significant molecular reprogramming during regeneration, whereas newly produced MPP2 and MPP3 are able to maintain a strong Meg and GM potential, respectively, hence highlighting the stability of their intrinsic lineage priming. Altogether, they identify MPP2/MPP3 as a compartment of myeloid amplification emerging rapidly from regenerating HSC^{LT}, and suggest that the molecular reprogramming of MPP4 also likely contributes to the initial burst of myeloid cell production.

Functional Reprogramming in Regenerating HSCs

To further investigate the changes in self-renewal activity occurring in regenerating HSC^{LT}, we re-transplanted donor-derived HSC^{LT} isolated at 2 and 3 weeks post-transplantation into lethally irradiated secondary recipients and compared their engraftment to freshly isolated control HSC^{LT} (Figure 7A).

Strikingly, both 2 and 3 week regenerating HSC^{LT} showed a nearly complete lack of reconstitution activity in PB output and BM engraftment compared to control HSC^{LT} (Figures 7B, 7C, and S7A). Limit dilution analyses performed with unfractionated BM cells directly confirmed the impaired engraftment ability of 2 week regenerating HSC^{LT} independently of surface markers (Figures 7D and S7B). Both 2 and 3 week regenerating HSC^{LT} also clearly displayed an activated status, characterized by decreased expression of the quiescence-enforcing transcription factors *Foxo3a* and *Egr1*, increased expression of *Slamf1* and the MPP markers *Cd48* and *Flt3*, and induction of the cell-cycle machinery with increased expression of G2-to-M phase cyclin/CDK complex genes (Figure 7E). Short-term in vivo 5-ethynyl-2'-deoxyuridine (EdU) incorporation experiments directly confirmed the highly proliferative status of 2 week regenerating HSC^{LT}, which had essentially returned to steady-state levels by 3 weeks (Figure 7F). In fact, by 4 weeks post-transplantation, donor-derived HSC^{LT} had also restored normal engraftment ability in vivo and plating activity in methylcellulose (Figures S7C and S7D). Finally, cytokine array analyses showed minimal perturbation in cytokine levels (including M-CSF and IL-1) except for increased IL-6 production in the BM cavity of transplanted mice (Figures 7G and S7E; data not shown). Taken together, these results demonstrate that regenerating HSC^{LT} temporarily lose their self-renewal activity as they overproduce myeloid-biased MPPs, and only recover their normal engraftment when blood homeostasis and adequate production of lymphoid-primed MPP4 are restored.

DISCUSSION

Here, we show that the MPP compartment is composed of at least two distinct subsets of myeloid-biased MPPs (i.e., MPP2 and MPP3), which work together with lymphoid-primed MPP4 to regulate blood production. We find that all three subsets of lineage-biased MPPs are independently produced by HSCs, but with different kinetics and at varying levels depending on hematopoietic demands. We demonstrate that myeloid-biased MPPs are important for maintaining blood homeostasis at steady state, and for rebuilding the myeloid lineages in regenerative conditions. Taken together, our results support a model of blood development that highlights the key role of the MPP compartment, and places its various lineage-biased subsets at the center stage of lineage specification. They also support the idea that the HSC compartment is functionally plastic, with HSCs committed to differentiation and supporting blood regeneration temporarily losing their engraftment ability. This updated scheme has important implications for understanding HSC self-renewal and blood production in various physiological contexts.

Competing Models of Blood Development

HSCs are extremely efficient at sustaining long-term multilineage hematopoietic reconstitution in vivo and are the only blood cells amenable to single cell transplantation assays. In contrast, the functional characterization of multipotent or lineage-committed progenitors remains challenging due to their low yield and transient output in transplantation assays, which pre-

clude most clonal analyses except for the recent attempt at lentiviral barcoding of MPP4 (Naik et al., 2013). Consequently, current models of blood development place significantly more emphasis on HSC function as measured by transplantation rather than on the activity of non-self-renewing MPPs to explain lineage specification. In the “clonal composition” model, the distinct patterns of blood reconstitution in single cell transplantation assays are explained by selection of particular HSC clones (Copley et al., 2012), and the myeloid bias associated with old age by a shift in clonal dominance with over-representation of myeloid-biased HSCs (Beerman et al., 2010). Consistent with this idea, myeloid-biased HSCs have been prospectively identified based on Hoechst dye exclusion (Challen et al., 2010) or high CD150 expression (Morita et al., 2010). Both the “graded differentiation” and “myeloid bypass” models directly argue against the existence of true MPPs and propose instead either a gradual loss of lineage potential with differentiation, starting with loss of Meg potential at the HSC stage (Adolfsson et al., 2005) and dendritic cell (DC) potential at the MPP4 stage (Naik et al., 2013), or the existence of long-lived myeloid-restricted repopulating progenitors that produce myeloid cells independently from other lineages (Yamamoto et al., 2013). However, all these models are interpretations of reconstitution patterns in transplanted mice, which are unlikely to reflect HSC function and lineage specification in native conditions as directly suggested by two recent lineage tracking studies that both showed a limited contribution of HSCs to ongoing hematopoiesis (Sun et al., 2014; Busch et al., 2015). It is also clear that engraftment and quiescence are intrinsically connected, and that more metabolically active MPP1 are less engrafting than quiescent HSCs (Cabezas-Wallscheid et al., 2014). However, it remains unclear whether MPP1 are truly separable from HSCs or whether they represent a reversible stage of HSC activation. This unfortunately cannot be directly tested using transplantation approaches.

To reconcile these observations, we propose a “dynamic model” of blood development in which the HSC compartment is, in fact, composed of distinct subsets of quiescent, activated, and lineage-primed HSCs that represent a continuum of likely reversible states and encompass the currently described HSC^{LT} (HSC/MPP1) and HSCST/ITRC populations. In turn, this plastic HSC compartment conveys lineage specification through the independent production of distinct subsets of lineage-biased MPPs in proportions that are based on hematopoietic demands (Figure 7H). This differs substantially from the prevailing views that consider HSCs solely as an engrafting population rather than a continuum of functional activity, and MPPs as a single linear progression of differentiation rather than parallel pathways that independently control the output of the myeloid and lymphoid lineages. The capacity of HSCs to independently produce specific subsets of lineage-biased MPPs is likely to contribute to the clonal heterogeneity observed upon single cell transplantation. Hence, myeloid-biased HSCs and long-lived myeloid-restricted repopulating progenitors could reflect the reconstitution pattern of HSCs that are primed to produce myeloid-biased MPP2 and MPP3, with balanced HSCs reflecting the behavior of uncommitted HSCs and lymphoid-biased HSCs of HSCs that are poised to produce lymphoid-primed MPP4.

Parallel MPP Compartments

Our results considerably expand on the functional characterization of the MPP compartment. We show that while fully multipotent in permissive culture conditions, both myeloid-biased MPPs display specific lineage preferences (i.e., Meg for MPP2 and GM for MPP3). We confirm these biases at the molecular level, with lineage priming reflective of the dominant blood output for each MPP subset, but otherwise only modest differences in their global transcriptional networks as expected for multipotent progenitors. In the case of the lymphoid-primed MPP4, it is also likely that their strong GM poising directly contributes to their ability to produce myeloid cells at steady state and to be quickly reprogrammed toward exclusive myeloid output in regenerative conditions. We validate these lineage biases *in vivo* by showing that MPP2 and MPP3 are more efficient at producing GMPs than MPP4, and by establishing the importance of MPP2 for megakaryopoiesis using *Mpl*^{-/-} mice. Moreover, we show that HSCs independently produce all MPP subsets, but at different levels and with different kinetics as exemplified by the delayed emergence of the lymphoid-primed MPP4 in regenerative conditions and the faster production of the Meg-biased MPP2 *in vitro*. This is interesting in the context of the strong Meg potential described for HSCs further purified with Von Willebrand factor (Sanjuan-Pla et al., 2013), which could directly reflect *in vivo* enrichment of HSCs that are primed to produce Meg-biased MPP2. Our results also indicate that Flk2⁻ myeloid-biased MPPs are produced independently of Flk2⁺ MPP4, which contradicts recent fate-mapping results suggesting that all blood cells are derived from Flk2-expressing progenitors (Boyer et al., 2011). However, because MPP2 and MPP3 both express *Flk2* mRNA at low levels at steady state and at increasing levels during regeneration, it is possible that the transgenic *Flk2-Cre* used in these experiments is activated in these MPP subsets despite their lack of Flk2 surface protein expression. Another possibility is that the early window during which MPP2 and MPP3 rebuild the myeloid lineage, before re-establishment of homeostasis and the dominant contribution of Flk2⁺ MPP4, has been missed in this study. It will therefore be important to use additional *in vivo* tracking approaches, such as those recently published in native conditions (Sun et al., 2014; Busch et al., 2015), to precisely map the differentiation paths of these MPP subsets.

Molecular Reprogramming in Regenerative Conditions

Consistent with their lineage biases and specific roles in myeloid cell production, we show that the lineage programming of MPP2 and MPP3 remains largely unchanged during regeneration. In contrast, HSCs and MPP4 show transient reprogramming of their lineage fate, with altered expression of similar myeloid lineage genes during blood regeneration. This is characterized by downregulation of *Cebpa/Irf8* and concomitant upregulation of *Runx1/Pu.1*, suggesting that the rebalancing of these two sets of transcription factors plays an important role in priming myeloid lineage output in regenerative conditions. Indeed, C/EBP α is dispensable for regenerative “emergency” granulopoiesis (Hirai et al., 2006), and its downregulation may represent a mechanism to activate HSC proliferation (Ye et al., 2013). Recent evidence also suggests that elevated levels of Pu.1, a direct target of Runx1, are crucial for promoting rapid myeloid differentiation in

HSCs (Mossadegh-Keller et al., 2013). These reciprocal changes could therefore enforce HSC proliferation and transiently favor HSC differentiation along a myeloid-biased MPP pathway, with decreased expression of *Irf8* eventually limiting monocyte/DC specification in the context of elevated Pu.1 activity (Tamura and Ozato, 2002). The transient induction of this program in MPP4 could directly enforce myeloid output in a population that, despite its GM priming, is also beginning to upregulate key lymphoid differentiation genes. The increased IL-6 levels detected in the BM cavity of transplanted mice could be part of this mechanism as IL-6 is known to reprogram MPP4 away from lymphoid differentiation and to amplify myeloid differentiation (Reynaud et al., 2011).

Implications for Blood Production

Our results support the idea that myeloid-biased MPPs serve as a transient compartment of myeloid expansion that can be rapidly activated by HSCs to ensure appropriate production of myeloid cells (Figure 7H). They also demonstrate that regenerating HSCs temporarily lose self-renewal activity as they overproduce myeloid-biased MPPs and rebuild the blood system, a phenotype perhaps linked to their increased cell cycle activity, activated metabolic status, and/or myeloid priming (Kohli and Passequé, 2014). This finding raises further caution about defining HSC identity solely based on transplantation experiments because regenerating HSCs do not perform well in this assay despite being fully capable of maintaining blood production over time. It also raises the interesting possibility that increased CD150 expression observed in both poorly engrafting myeloid-biased HSC clones (Morita et al., 2010; Beerman et al., 2010) and regenerating HSCs is linked to the production of myeloid-biased MPPs and is essentially a reflection of their activated/primed status. It will now be interesting to explore whether differential production of lineage-biased MPPs by HSCs could account for other long-lasting changes in blood production, such as the predominant lymphoid/GM output of fetal hematopoiesis (Kawamoto, 2006); the GM skewing, immunosenescence, and anemia of the old blood system (Geiger and Rudolph, 2009); or the aberrant overproduction of different myeloid lineages in hematological malignancies (Tefferi and Gilliland, 2007). It will also be exciting to determine whether similar populations of myeloid-biased MPPs exist in humans. In addition, one of the main complications of anti-cancer therapies or BM transplantation protocols is their deleterious effect on the blood system, leading to prolonged neutropenia and increased risk for infections. It will therefore be compelling to test whether HSC differentiation pathways can be manipulated to favor production of specific lineage-biased MPPs and thereby optimize blood recovery following hematopoietic injury or rebalance lineage output in an aging or deregulated blood system.

EXPERIMENTAL PROCEDURES

Mice and Flow Cytometry

Six- to 8-week-old CD45.2 C57Bl/6 wild-type or β -actin-*Gfp* (Forsberg et al., 2006) mice were used as donors for cell isolation, and 8- to 12-week-old CD45.1 C57Bl/6-Boy/J wild-type mice were used as recipients for cell transplantation. Transplanted mice were given antibiotic-containing water for 4 weeks. All mice were maintained at the University of California San

Francisco in accordance with Institutional Animal Care and Use Committee-approved protocols. Staining and enrichment procedures for flow cytometry were performed as previously described (Santaguida et al., 2009; Reynaud et al., 2011). Cells were sorted on a fluorescence-activated cell sorting (FACS) ARIALL and analyzed on an LSRII (Becton Dickinson) upon PI exclusion of dead cells. Each population was double sorted to ensure maximum purity.

In Vitro Analyses

Proliferation, BrdU incorporation, differentiation and OP9/OP9-DL1 co-culture experiments, microarray analyses, and Fluidigm experiments were performed as described in the Supplemental Experimental Procedures. Three to five independent biological replicates were used for each population. Methylcellulose and MegaCult assays were performed according to the manufacturer's protocols (Stem Cell Technologies).

In Vivo Analyses

Transplantations, lineage tracking, and CFU-S assay were performed as previously described (Forsberg et al., 2006; Reynaud et al., 2011). Congenic recipient mice (CD45.1) were either lethally (1,100 rad, split dose 3 hr apart) or sub-lethally (950 rad, split dose 3 hr apart) irradiated. Purified donor cells were injected into the retro-orbital plexus, and hematopoietic reconstitution was monitored over time in the peripheral blood based on CD45.2 or GFP expression. EdU incorporation experiments and cytokine analyses were performed according to the manufacturer's protocols (Life Technologies).

Statistics

All data are expressed as mean \pm SD or SEM as indicated. The p values were generated using unpaired Student's t test or a Mann-Whitney u test (Fluidigm) and considered significant when ≤ 0.05 . N indicates the number of independent experiments performed.

ACCESSION NUMBERS

The accession number for the microarray data reported in this paper is GEO: GSE68529.

SUPPLEMENTAL INFORMATION

Supplemental Information includes Supplemental Experimental Procedures, seven figures, and four tables and can be found with this article online at <http://dx.doi.org/10.1016/j.stem.2015.05.003>.

AUTHOR CONTRIBUTIONS

E.P. conceptualized the experiments; E.M.P., D.R., and E.P. designed the methodology; E.M.P., D.R., and Y-A.K. performed the investigation; D.C. and J.M.S. performed microarray analysis; F.J.C-N. and B.G. performed the Fluidigm analyses; A.D.L. provided resources for the study; E.M.P., D.R., and E.P. wrote the original draft; E.M.P., Y-A.K., and E.P. reviewed and edited the manuscript; and E.M.P., B.G., and E.P. obtained funding for the study.

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Cell Stem Cell

Supplemental Information

Functionally Distinct Subsets of Lineage-Biased Multipotent Progenitors Control Blood Production in Normal and Regenerative Conditions

Eric M. Pietras, Damien Reynaud, Yoon-A Kang, Daniel Carlin, Fernando J. Calero-Nieto, Andrew D. Leavitt, Joshua M. Stuart, Berthold Göttgens, and Emmanuelle Passegué



Figure S1, Functional characterization of BM LSK subsets, related to Figure 2. (A) Myeloid differentiation potential in liquid culture. Representative FACS plots showing the kinetics of FcγR and Mac-1 acquisition in the indicated populations (n ≥ 2). (B) B cell differentiation on OP9/IL-7 stromal cells. Representative FACS plots showing the kinetics of CD19⁺ B-cell and Mac-1⁺ myeloid (M) cells differentiation over time. Results are representative of 4 independent experiments. (C) Representative FACS plots showing unique B and mix B/M output from single MPP3 and MPP4 cultured for 16 days on OP9/IL-7 stromal cells. The frequency for each output is indicated. (D) T cell differentiation on OP9-DL1 stromal cells. Representative FACS plots showing the kinetics of CD25 and CD44 acquisition in the indicated populations. Results are representative of 4 independent experiments. (E) Representative FACS plots showing that MPP2 and MPP3 are phenotypically related to Pre-MegE and Pre-GM progenitors, respectively.



Figure S2, *In vivo* reconstitution assays, related to Figure 3. (A) CD45.2 HSC^{LT} or HSCST were transplanted into lethally irradiated CD45.1 recipients (50 cells/mice) together with 3×10^5 Sca-1-depleted CD45.1 BM cells. The table and FACS plots illustrate engraftment and percentage of CD45.2⁺ donor-derived myeloid (Mac-1⁺) and lymphoid (B220⁺/CD3⁺) cells in the peripheral blood at 4 and 16 weeks post-transplantation. Results are expressed as mean ± SD. (B) Representative FACS plots showing examples of nucleated myeloid cells (Mac-1⁺/Gr-1⁺), B cells (B220⁺) and T cells (CD3⁺) produced by the indicated donor GFP⁺ populations 27 days post-transplantation. (C) Representative FACS plots showing examples of platelets (Ter-119⁺/Gr-1⁺/CD61⁺) produced by the indicated donor GFP⁺ populations 20 days post-transplantation.

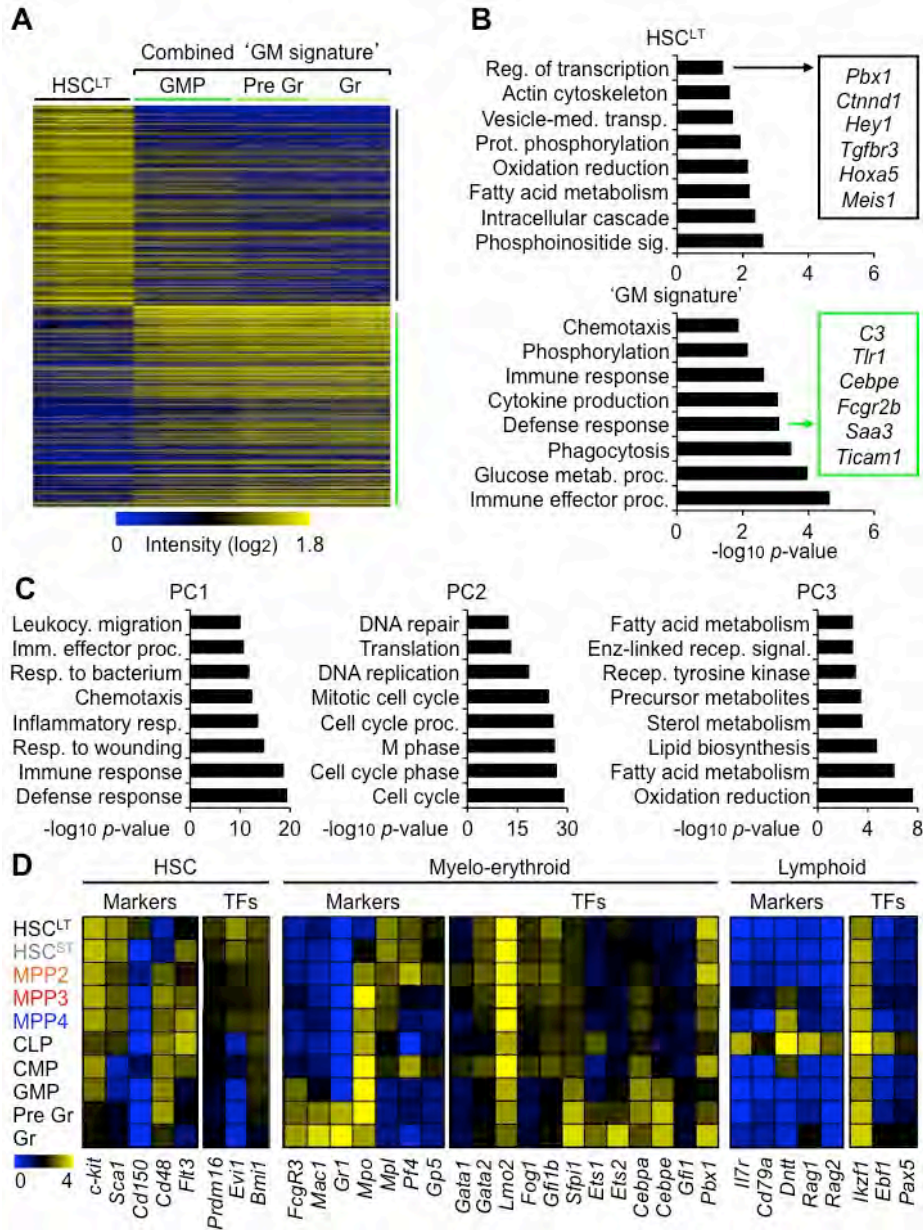


Figure S3, Molecular characteristics of lineage-biased MPP subsets, related to Figure 4. (A) Individual gene signatures representing the 1000 most enriched genes expressed in HSC^{LT} and GM lineage-committed cells. GMP: Lin⁻/c-Kit⁺/Sca-1⁻/CD34⁺/FcγR⁺; Pre Gr: Mac-1⁺/Gr-1^{int}; Gr: Mac-1⁺/Gr-1⁺. (B) GO analyses of the gene signatures shown in (A). (C) GO analyses of the PC components shown in Figure 4B. (D) Heat map showing normalized dye intensity for expression of key hematopoietic markers (markers) and transcription factors (TFs) in the indicated populations.

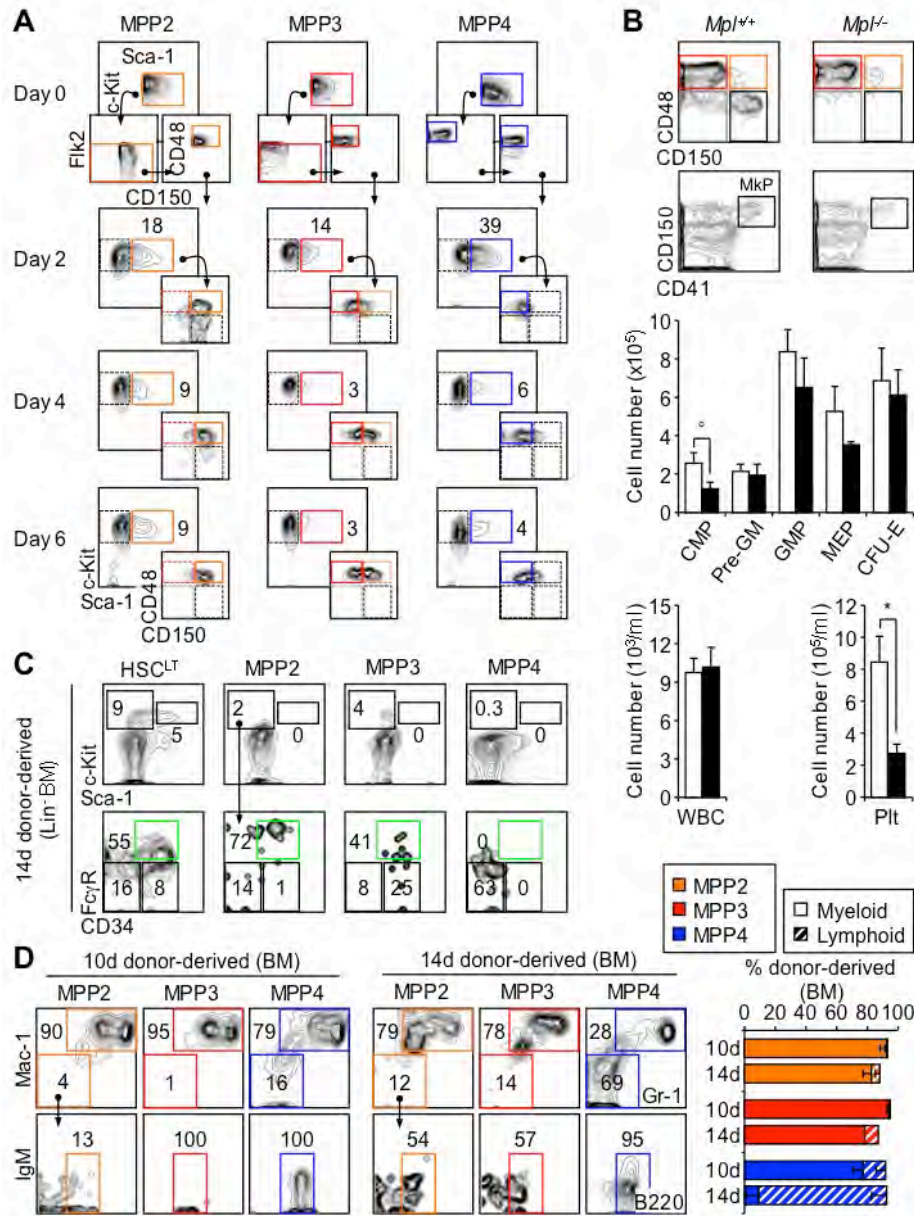


Figure S4, Hierarchy of lineage-biased MPPs, related to Figure 5. (A) Differentiation *in vitro*. Representative FACS plots showing kinetics of MPP2, MPP3 and MPP4 differentiation under myeloid culture conditions. Results are representative of 2 independent experiments. (B) Representative FACS plots and size of the indicated stem and progenitor populations in the BM and blood of *Mpl*^{+/+} and *Mpl*^{-/-} littermates (3 mice/group). (C) Differentiation *in vivo*. Representative FACS plots of LSK and myeloid progenitor output 14 days following transplantation of 5,000 HSC^{LT}, MPP2, MPP3 and MPP4 (3 mice/group). (D) Representative FACS plots (left) and quantification (right) of donor-derived myeloid and lymphoid BM output from the indicated populations 10 and 14 days after transplantation (3-5 mice/group). Results are mean ± SD. °p < 0.05, *p < 0.001.

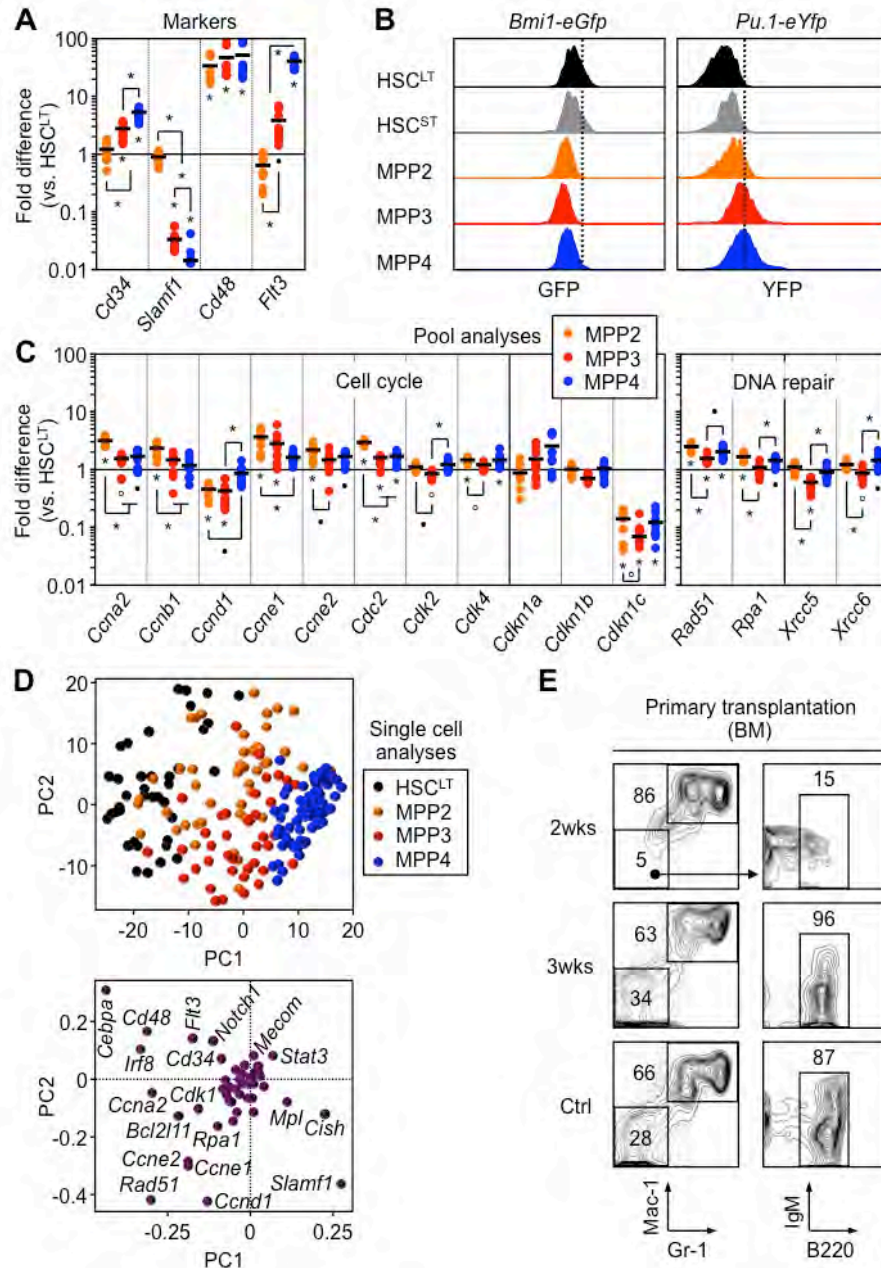


Figure S5, Molecular regulation at steady state, related to Figures 5 and 6. (A) Fluidigm gene expression analyses of surface markers. Results are expressed as mean (bar) and individual fold differences compared to HSC^{LT} (8-12 pools of 100 cells/). (B) GFP expression in the indicated BM LSK subsets of *Bmi1-eGfp* and *Pu.1-eYfp* reporter mice. (C) Fluidigm gene expression analyses of cell cycle and DNA repair genes. Results are expressed as in (A). (D) PC analysis (top) and loading association (bottom) of Fluidigm gene expression data acquired from single cells (30-58 cells/population). (E) Representative FACS plots showing myeloid and lymphoid output in transplanted recipients at 2 and 3 weeks post injection compared to steady state control (Ctrl) mice. $^{\circ}p < 0.05$, $^{\bullet}p < 0.01$, $^*p < 0.001$.

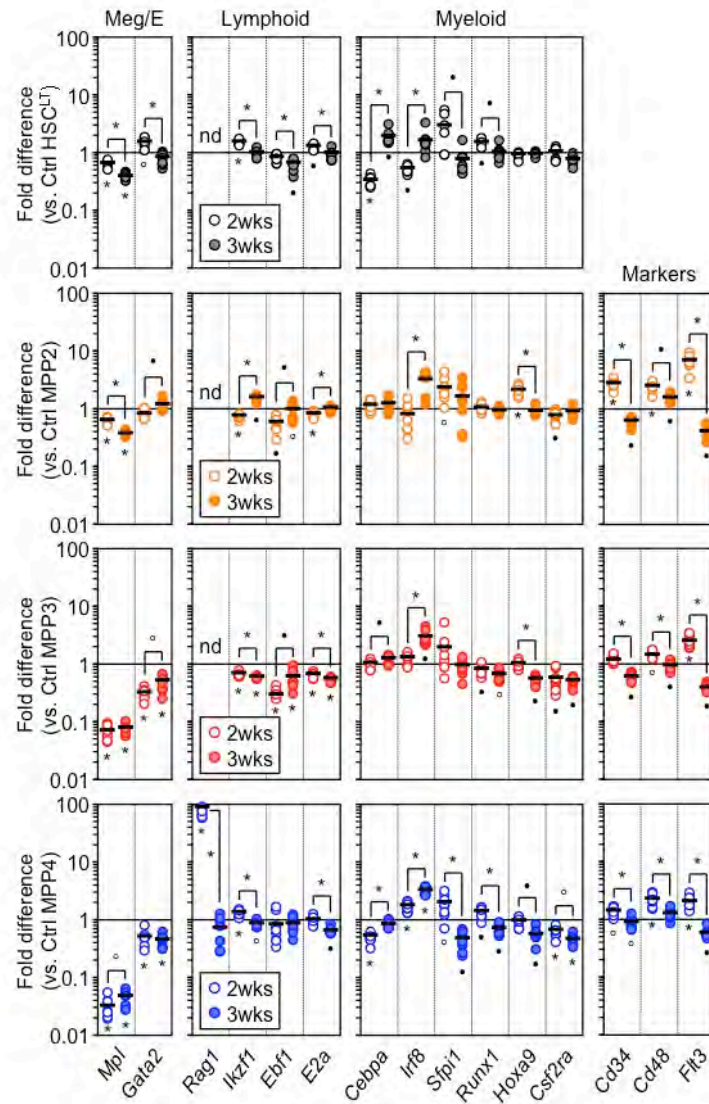


Figure S6, Molecular regulation during regeneration, related to Figure 6. Fluidigm gene expression analyses of key lineage determinant and surface marker genes in regenerating HSC^{LT} and newly produced MPP2, MPP3 and MPP4 at 2 and 3 weeks post-transplantation. Results are expressed as mean (bar) and individual fold differences compared to their respective steady state control (Ctrl) populations (8-12 pools of 100 cells/population). °p < 0.05, • p < 0.01, *p < 0.001.

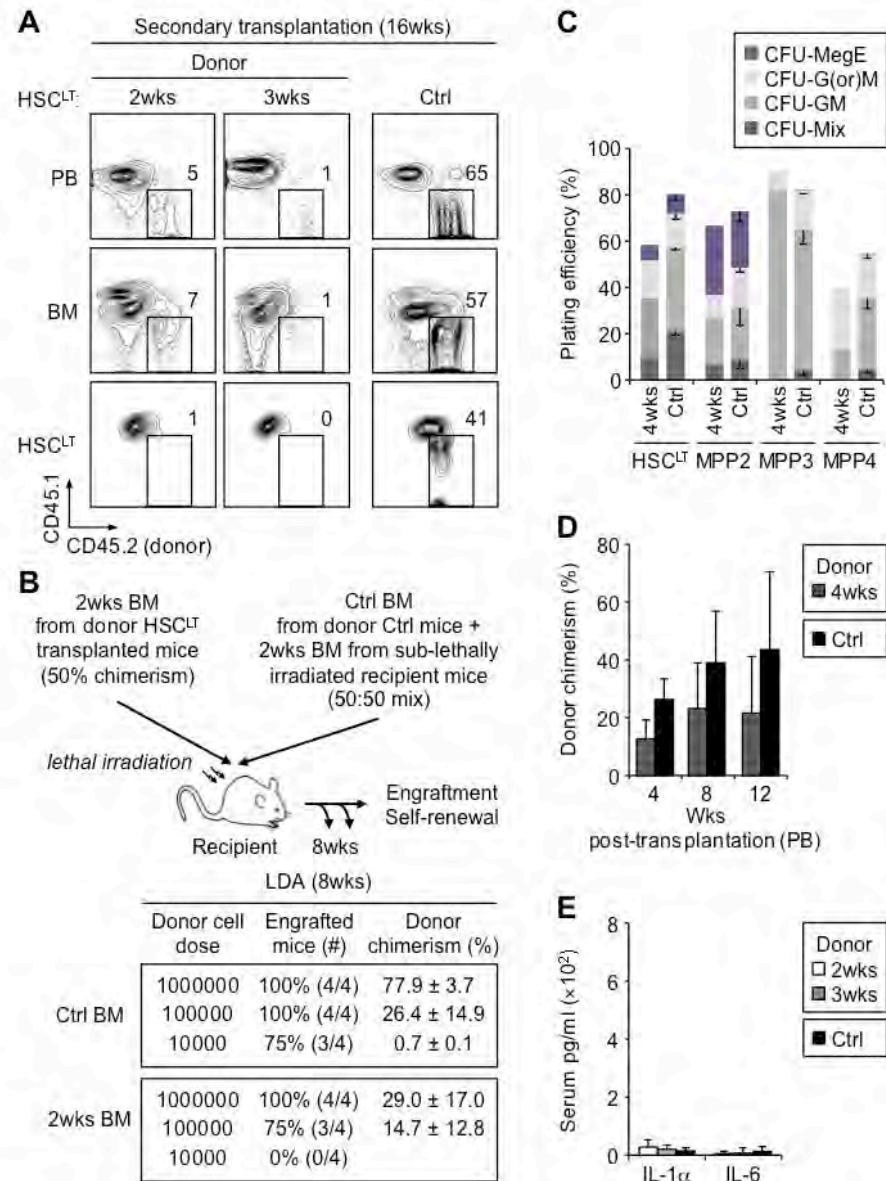


Figure S7, Regenerating hematopoiesis, related to Figure 7. (A) Representative FACS plots of CD45.2 donor chimerism in secondary recipients transplanted with 2 and 3 weeks regenerating HSC^{LT} and control (Ctrl) HSC^{LT}. (B) LDA experimental scheme (top) and detailed results at 8 weeks post-transplantation (bottom). Lethally irradiated CD45.1 recipients were transplanted with various doses of BM cells obtained either from mice transplanted with donor CD45.2 HSC^{LT} and re-isolated 2 weeks after transplantation (2wks BM; 50% chimerism) or from steady state control CD45.2 mice and mixed 1:1 with BM cells isolated from CD45.1 recipient 2 weeks after sub-lethal irradiation (Ctrl BM; 50:50 mix). Transplanted mice were bled every month and analyzed for engraftment and donor chimerism. Mice with $\geq 0.5\%$ donor PB chimerism at 8 weeks post-transplantation were considered engrafted. (C) Methylcellulose clonogenic assays of 4 weeks regenerating HSC^{LT}, MPP2, MPP3 and MPP4. Ctrl are the same as in Figure 6D. Results are expressed as mean \pm SEM (n = 1-4). (D) Engraftment over time of 4

week regenerative HSC^{LT} in PB. Ctrl are the same as in Figure 7B. Results are expressed as mean \pm SD (5-13 mice/group). (E) IL-1 and IL-6 levels in PB serum. Results are expressed as mean \pm SD (9-12 mice/group).

Table S1, Gene signatures and GO analysis of HSC^{LT} and GM lineage-committed populations, related to Figure 4. List of top 1000 most highly expressed genes in each population relative to the indicated reference populations, ranked based on SAM score. *p*- and *q*-values for the 1000th probe in each signature are displayed at the top of the list. GO categories are listed with accession number, enrichment *p*-value, and genes contributing to the enrichment. The highlighted GO categories are shown in Figure S3A.

Table S2, GO analysis of genes driving PC separation, related to Figure 4. Complete list of GO categories enriched in the top 1000 genes driving each principal component. The table displays accession number, enrichment *p*-value, and genes contributing to the enrichment. The highlighted GO categories are shown in Figure S3C.

Table S3, Gene signatures and GO analysis of MPP2, MPP3 and MPP4, related to Figure 4. List of top 1000 most highly expressed genes in each population relative to the indicated reference populations, ranked based on SAM score. *p*- and *q*-values for the 1000th probe in each signature are displayed at the top of the list. GO categories are listed with accession number, enrichment *p*-value, and genes contributing to the enrichment. The highlighted GO categories are shown in Figure 4D.

Table S4, Overlap with published MPP gene expression datasets, related to Figure 4. Complete list of genes overlapping between the 1000 most highly expressed genes in MPP2, MPP3 and MPP4, and published microarray datasets. Genes are ranked based on SAM score. Dataset accessions are GSE55525, GSE15907 or as described (Cabezas-Wallscheid et al., 2014).

Supplemental Experimental Procedures

Mice

Six- to 8-week-old CD45.2 C57Bl/6 wild type or *β-actin-Gfp* (Forsberg et al., 2006) mice were used as donor for cell isolation, and 8-12 week old CD45.1 C57Bl/6-Boy/J wild type mice as recipients for cell transplantation. Transplanted mice were kept on antibiotic-containing water for 4 weeks. *Bmi1-Gfp* (Hosen et al., 2007), *Pu.1-eYFP* (Kirstetter et al., 2006) and *Mpl^{-/-}* (Gurney et al., 1994) mice were used for hematopoietic analyses. Respective littermates were used as controls. All mice were maintained at UCSF in accordance with IACUC approved protocols.

Flow cytometry

Staining and enrichment procedures for stem and progenitor cells were performed as previously described (Santaguida et al., 2009; Reynaud et al., 2011). Briefly, unfractionated or c-Kit-enriched BM cells or splenocytes were stained with unconjugated rat anti-lineage (B220, CD3, CD4, CD5, CD8, Ter119, Mac-1, Gr-1, IL-7Ra) antibodies followed by goat anti-rat-Cy5PE or Qdot605 secondary antibodies. Cells were then stained with directly conjugated c-Kit-APC-eFluor780, Sca-1-PB, Flk2-bio/streptavidin-Cy7PE, CD48-A647, CD150-PE, CD34-FITC and FcγR-PerCP-eFluor710 antibodies. For particular myeloid progenitor analyses (Pronk et al., 2007), unfractionated BM cells were stained with unconjugated rat anti-lineage antibodies followed by goat anti-rat-Cy5PE antibodies and directly conjugated c-Kit-APC-eFluor780, Sca-1-PB, CD150-APC, CD41-PE, CD105-FITC and FcγR-PerCP-eFluor710 antibodies. For additional surface marker analyses, unfractionated BM cells were stained for lineage markers as above and subsequently stained with directly conjugated c-Kit-APC-eFluor780, Sca-1-PB, Flk2-bio/streptavidin-Cy7PE, CD48-A647, CD150-PE (LSK populations) or c-Kit-APC-eFluor780, Sca-1-PB, CD34-Bio/streptavidin-Cy7PE and FcγR-PerCP-eFluor710 (myeloid progenitors) plus ESAM-FITC or CD41-FITC. Blood and BM mature cells were stained with directly conjugated B220-APC-A750, Gr-1-PB, Mac-1-Cy7PE, CD3-APC and Ter-119-Cy5PE antibodies supplemented with CD45.2-FITC and CD45.1-PE antibodies for transplantation studies. For analysis of donor-derived stem and progenitor populations in transplanted mice, unfractionated BM cells were stained for lineage markers as above and subsequently stained with directly conjugated c-Kit-APC-eFluor780, Sca-1-PB, Flk2-bio/streptavidin-Qdot605, CD48-A647, CD150-PE (LSK populations) or c-Kit-APC-eFluor780, Sca-1-PB, CD34-Bio/streptavidin-Qdot 605 and FcγR-PerCP-eFluor710 (myeloid progenitors), plus CD45.2-FITC and CD45.1-Cy7PE antibodies. Cells were sorted on a FACS ARII and analyzed on an LSRII (Becton Dickinson) upon PI exclusion of dead cells. Each population was double sorted to ensure maximum purity.

In vitro assays

For expansion in liquid culture, cells (500 cells/well in 96-well plates) were grown in IMDM (Invitrogen) supplemented with 5% FBS (StemCell Technology), L-glutamine (2 mM), penicillin (50 U/ml)/streptomycin (50 µg/ml) and the following cytokines (all from PeproTech): SCF (25 ng/ml), Flt3L (25 ng/ml), IL-11 (25 ng/ml), IL-3 (10 ng/ml), GM-CSF (10 ng/ml), TPO (25 ng/ml) and EPO (4 U/ml). Cell numbers were determined by trypan blue exclusion of dead cells using a hemacytometer. For BrdU incorporation experiments, cells (5×10^3 to 10^4 cells/well in

96-well plates) were cultured for 1 hour in the presence of 60 μ M BrdU (Sigma), fixed in PBS/4% paraformaldehyde and permeabilized in PBS/0.2% Triton-X100. After DNase I treatment, cells were incubated with FITC-conjugated anti-BrdU antibody (Pharmingen), washed and resuspended in PBS/0.2% for analysis as described (Santaguida et al., 2009). For methylcellulose assays, cells (1 cell/100 μ l/well in 96-well plates or 100 cells/1 ml in 3 cm dish) were cultured in M3231 methylcellulose (Stem Cell Technologies) supplemented as for the liquid cultures. Colonies were counted after 10 days of culture, and for the clonogenic assays at least 60 wells were monitored per cell type for each experiment. For MegaCult assays, cells ($250\text{--}1 \times 10^3$ cells/1 ml in 3 cm dish) were plated and grown for 5 days in collagen-based media (StemCell Technologies) supplemented with TPO (50 ng/ml), IL-6 (20 ng/ml), IL-11 (50 ng/ml) and IL-3 (10 ng/ml). After fixation, Meg colonies were revealed by immunohistochemistry using an anti-CD41 antibody and scored upon visual inspection. For myeloid differentiation and surface marker analyses, cells (1,000 to 3,000 cells/well in 96-well plates) grown in StemPro media (Life Technologies) supplemented with penicillin (50 U/ml)/streptomycin (50 μ g/ml), SCF (25 ng/ml), IL-11 (25 ng/ml), Flt3L (25 ng/ml), IL-3 (10 ng/ml) and GM-CSF (10 ng/ml). Cells were analyzed after different culture periods by flow cytometry using c-Kit-APC-A750, Sca-1-PB, CD150-PE, CD48-APC, Mac-1-FITC and Fc γ R-Cy5.5PE antibodies. For lymphoid differentiation assays, cells (1 to 500 cells/well in 96-well plates) were sorted into wells pre-seeded with 5×10^5 OP9 (B cells) and OP9-DL1 (T cells) stromal cells as previously described (Reynaud et al., 2008). For B cell differentiation, cells were cultured in OptiMEM (Invitrogen) supplemented with 5% FBS in presence of SCF (10 ng/ml), Flt3L (10 ng/ml) and IL-7 (5 ng/ml). After sequential withdrawal of Flt3L and SCF, cells were maintained with IL-7 and analyzed after different culture periods by flow cytometry using CD19-PB and Mac-1-APC antibodies. For T cell differentiation, cells were cultured in OptiMEM supplemented with 5% FBS in presence Flt3L (10 ng/ml) and IL-7 (5 ng/ml), and analyzed by flow cytometry using CD25 biotin/Streptavidin-Cy7PE and CD44-APC antibodies.

***In vivo* assays**

Transplantations and lineage tracking experiments were performed as previously described (Forsberg et al., 2006; Reynaud et al., 2011). Congenic recipient mice (CD45.1) were either lethally (1100 rad, split dose 3h apart) or sub-lethally (950 rad, split dose 3h apart) irradiated using a cesium source. Purified donor cells were injected into the retro-orbital plexus and hematopoietic reconstitution was monitored over time in the peripheral blood based on CD45.2 or GFP expression. For sub-lethal irradiation, purified donor cells alone were injected and transplanted mice were first bled after 7 days and then every 3-5 days for up to 34 days. For lethal irradiation, unfractionated donor BM cells alone, or purified donor cells together with 300,000 Sca-1-depleted congenic helper BM cells, were injected and transplanted mice were first bled after 4 weeks and then on a monthly schedule. Peripheral blood was obtained from retro-orbital bleeding, and either collected in 4 ml of ACK (150 mM NH_4Cl /10 mM KHCO_3) containing 10 mM EDTA for flow cytometry analyses, or in EDTA-coated Microtainer tubes (Beckton-Dickenson) for complete blood counts using an Hemavet hematology system (Drew Scientific). Results of limit dilution analyses (LDA) were calculated with the ELDA software (<http://bioinf.wehi.edu.au/software/elda/>). Spleen colony-forming unit (CFU-S) assays were performed as described (Forsberg et al., 2006). Lethally irradiated recipient mice were transplanted with 50 to 250 cells of the indicated populations. Spleens were harvested 12 days

after transplantation and fixed in Tellyesniczky's solution to determine CFU-S₁₂ frequency (numbers of injected cells divided by numbers of observed spleen colonies). For EdU incorporation experiments, mice were injected with 10 mM EdU (Life Technologies) 1h prior harvest. Purified cells were pipetted onto poly-lysine coated slides (500-2,000 cells per slide) and EdU incorporation was detected using A594-labeled azide click chemistry according to the manufacturer's instruction (Life Technologies). For Wright-Giemsa staining, purified cells were cytopspun onto glass slides (1,000-5,000 cells per slide) and processed according to standard protocols.

Cytokine analyses

For PB serum, blood was harvested from euthanized mice via cardiac puncture, allowed to coagulate at room temperature for 30 min, and subsequently spun down at 12,000 × g for 10 min to remove blood cells. For BM fluid, the four long bones (two femurs and two tibiae) of the same mice were flushed with 150-200 µl HBSS/2% FBS using a 0.3cc insulin syringe with a 28g needle and spun at 500 × g for 5 min to remove BM cells. Supernatants were further clarified by spinning down at 12,000 × g for 10 min, and samples were subsequently stored at -20°C until use. For cytokine measurement, 50 µl of 2x-diluted sample was analyzed with a Luminex Cytokine Mouse 20-plex panel (Life Technologies) using a BioPlex instrument (Bio-Rad) according to the manufacturer's instructions.

Microarray analyses

For microarray analyses, three to five independent biological replicates were used for each population. Total RNA was isolated from 2x10⁴ purified cells directly sorted into TRIzol-LS (Invitrogen) and purified using Arcturus PicoPure (Applied Biosystems) with RNase-free DNase (Qiagen). RNA was amplified, labeled, and fragmented using NuGEN Ovation Pico linear amplification kits (Nugen Technologies) and hybridized onto mouse Gene ST 1.0 arrays (Affymetrix). Gene signatures were determined by contrasting the gene expression patterns characteristic of each population (MPP2, MPP3, MPP4, HSC^{LT} and GM lineage-committed cells) against the other indicated population(s). Gene expression microarray data were normalized using RMA followed by quantile normalization as implemented in the 2.15.1 R package (www.r-project.org) using a standard (lambda=1) exponential reference distribution. For a specific population a dichotomous division of the samples was defined by labeling the population (*e.g.*, MPP3) as the first class and the other two populations (*e.g.*, MPP2 and MPP4) as the second class. Significance Analysis of Microarrays (SAM) (Tusher et al., 2001) was then performed using the expression data and the dichotomous labeling of the data to determine SAM *delta* scores, which reflect the degree of differential expression of each gene in the first class compared to the second. SAM performs a robust version of a two-way test that safeguards the analysis against extremely low variance estimates associated with genes of low expression. The top 1000 most highly expressed genes were collected as the gene signature for each population. Reported p- and q-values for the 1000th probe in each gene signature were derived via modified t-test and false discovery rate (FDR) with 100 label permutations (Storey and Tibshirani, 2003). Microarray data were clustered by Pearson correlation with complete linkage using MultiExperiment Viewer 4.5.1. PC analysis of microarray data was performed using the prcomp function in R. Functional pathways representative of each gene signature were analyzed for

enrichment in gene categories from the Gene Ontology (GO) database (Gene Ontology Consortium, 2010) using DAVID Bioinformatics Resources ver. 6.7 (Huang et al., 2009). For gene signature and each GO category, the significance of the number of overlapping genes in the two sets was calculated using a Fisher's Exact test performed by the DAVID software. The P -value resulting from the test reflects the probability of obtaining the observed overlap or greater by chance. The $-\log_{10}$ of the P -value provides an intuitive level of significance and was used to rank GO categories. For principal component analysis the loadings of the centroids for each cell population in the first three principal components were used to generate plots. GO analyses were conducted using DAVID on the top 1000 genes contributing to each of the three principal components in its positive direction. Comparisons with MPP gene signatures from previously published microarray data were conducted either by running an independent SAM analysis on the published dataset to identify differentially expressed genes (Beerman et al., 2014) or, when fewer than three biological replicates were present per population (Gazit et al., 2013), differential expression was determined by subtracting one gene vector from the other. The top 1000 up or downregulated genes present in both datasets were then identified and overlapped with the gene signatures of MPP2, MPP3 and MPP4 described above. For RNAseq data (Cabezas-Wallscheid et al., 2014), differential expression results generated in this study were directly compared against the MPP2, MPP3 and MPP4 gene signatures described above.

Fluidigm analyses

Gene expression analyses using the Fluidigm 96.96 Dynamic Array IFC were performed as previously described (Reynaud et al., 2011). Briefly, pools of 100 cells or single cells were sorted directly into CellsDirect resuspension buffer (Invitrogen), reverse-transcribed and preamplified for 18 (bulk) or 22 cycles (single cells) using Superscript III Platinum Taq DNA polymerase (Invitrogen) using proprietary target-specific primers (Fluidigm). The resulting cDNAs were analyzed on a Biomark system (Fluidigm) using EvaGreen Sybr dye (Bio-Rad). Data were collected using Biomark Data Collection Software (Fluidigm). For bulk populations, data were analyzed using Biomark qPCR analysis software (Fluidigm) with a quality threshold of 0.65 and linear baseline correction, and identical settings used across all experiments performed on different 96.96 dynamic arrays. Melt curves and T_m values for each assay reaction were checked individually, and reactions with melt curves showing multiple peaks or poor quality (<0.65) were discarded, leaving 57 genes, excluding housekeeping genes (*Gapdh*, *Hprt*, *Tubb5* and *Gusb*) for further analysis. For gene expression quantification, data were exported as an Excel .csv file and analyzed by the $\Delta\Delta C_t$ method using *Gusb* for normalization as previously described (Reynaud et al., 2011). For PC analysis on bulk populations, data were subjected to quality control as described above and normalized to *Gapdh*. Housekeeping genes were removed prior to analysis, and genes with expression values above the range of the instrument ($C_t > 28$) were set to 29. PC analysis was then conducted using the `prcomp` function in R. For single cell gene expression analysis, 5 negative control wells (no cell sorted), 3 positive control wells (20 cells) and 88 wells containing a single cell were sorted for each population, corresponding to a single 96.96 dynamic array. Single-cell data were initially analyzed using Biomark qPCR analysis software as above. Of the 57 genes, those showing poor T_m quality in one or more reactions were removed from analysis, leaving 49 genes for further analysis (excluding housekeeping genes). Cells expressing fewer than 29 genes (poor quality) were also excluded, leaving 196 cells (40 HSC^{LT}, 45 MPP2, 41 MPP3, 70 MPP4) for further analysis. Expression

levels of *Gapdh* were used for normalization and PC analysis was conducted as above. tSNE analysis was performed using the MATLAB implementation (<http://homepage.tudelft.nl/19j49/t-SNE.html>) with standard settings (van der Maaten and Hinton, 2008). Data were preprocessed to keep only the first 28 components of PCA containing 95% of the information. Discriminant analysis classification was performed in Matlab using a quadratic classification.

Statistics

All the data are expressed as mean \pm standard deviation (SD) or standard error of the mean (SEM) as indicated. P values were generated using unpaired Student's t-test or a Mann-Whitney u-test (Fluidigm), and considered significant when ≤ 0.05 . N indicates the numbers of independent experiments performed.

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